The pace and prognosis of peripheral sensory loss in advanced age: association with gait speed and falls

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Abstract

Background: Peripheral sensory loss is considered one of many risk factors for gait impairments and falls in older adults, yet no prospective studies have examined changes in touch sensation in the foot over time and their relationship to mobility and falls. Therefore, we aimed to determine the prevalence and progression of peripheral sensory deficits in the feet of older adults, and whether sensory changes are associated with the slowing of gait and development of falls over 5 years.

Methods: Using baseline, and 18 and 60 month followup data from the Maintenance Of Balance, Independent Living, Intellect, and Zest in the Elderly (MOBILIZE) Study in Boston, MA, we determined changes in the ability to detect stimulation of the great toe with Semmes Weinstein monofilaments in 351 older adults. We used covariate-adjusted repeated measures analysis of variance to determine relationships between sensory changes and gait speed or fall rates.

Results: Subjects whose sensory function was consistently impaired over 5 years had a significantly steeper decline in gait speed (−0.23 m/s; 95% CI: -0.28 to −0.18) compared to those with consistently intact sensory function (−0.12 m/s; 95% CI: -0.15 to −0.08) and those progressing from intact to impaired sensory function (−0.13 m/s; −0.16 to −0.10). Compared to subjects with consistently intact sensation, those whose sensory function progressed to impairment during followup had the greatest risk of falls (adjusted risk ratio = 1.57 (95% confidence interval = 1.12 to 2.22).

Conclusions: Our longitudinal results indicate that a progressive decline in peripheral touch sensation is a risk factor for mobility impairment and falls in older adults.

Keywords: Neuropathy, Elderly, Mobility, Longitudinal, Feet

Background

Falls are a leading cause of morbidity and mortality among older adults, occurring in approximately one-third of community-dwelling persons over age 65 and costing over $31 billion in related injuries annually [1–3]. The presence of peripheral sensory loss is considered to be one of many different risk factors for falls [3–8]. A few previous studies demonstrated that older adults with peripheral sensory loss at the outset had an increased risk of subsequent falls [5–9]. However, these studies relied on single baseline measures of sensory function and did not examine the effect of changes over time [7, 8]. The longitudinal relationship between changes in peripheral sensory function and the development of falls is not known.

Peripheral neuropathy is common among older adults, [10] especially in those with diabetes [11, 12]. The prevalence of peripheral neuropathy has been reported to be approximately 7% in older adult populations worldwide [11, 13]. While there are many causes, including alcohol ingestion, vitamin B12 deficiency, cancer, chemotherapy, chronic kidney disease, and paraproteinemias, diabetes is the most frequent, accounting for 32–44% of patients with polyneuropathy [14, 15].
Many factors associated with peripheral neuropathy may increase the risk of falls, including impaired balance, muscle weakness, nutritional deficiencies, and medications [2, 16]. One mechanism by which the progression of sensory neuropathy may predispose people to falls is through alterations in gait. While slow gait speed has been reported to be cross-sectionally associated with distal sensory neuropathy [17], this could be either a cause of falls or protective mechanism to prevent falls. The present study takes advantage of longitudinal data from the Maintenance Of Balance, Independent Living, Intellect, and Zest in the Elderly (MOBILIZE) Study in Boston, MA, to assess the prevalence and progression of peripheral sensory deficits in the feet, and whether they are associated with slowing of gait and falls over a 5-year period.

Methods

Participants

The MOBILIZE Boston Study (MBS) is a prospective cohort study of a unique set of risk factors for falls in community-dwelling seniors living in the Boston area. The design and methodology for this study have been previously described in detail [18, 19]. In brief, 765 persons eligible for the MBS were enrolled using door-to-door population based recruitment. To be included, individuals had to be: > 70 years of age (or > 65 years if living with a participant), able to understand and communicate in English, able to walk 20 ft without personal assistance (walking aids permitted), expecting to live in the area for at least 2 years, and able to provide written informed consent. Exclusion criteria included terminal disease, severe vision or hearing deficits, and a Mini-Mental State Examination score < 18 [20, 21].

All subjects underwent a complete home and laboratory assessment of demographic characteristics, medical conditions, medications, functional status, gait speed, smoking status, alcohol use, blood pressure, and cerebral hemodynamics, then were followed prospectively for falls using a monthly postcard calendar (see below). These assessments were repeated at 18 months and 5 years, while monthly falls data were collected continuously.

The analysis described here utilized longitudinal data from three waves of MBS data collection over approximately five years. Of the original 765 MOBILIZE Boston participants, data from only 351 subjects were available for the current study because of death, institutionalization, or loss to followup. As expected, the study sample that survived the 5 year followup was healthier that the original cohort, with better scores on the Mini-mental State Examination [20], Trail-making Test [22], Short Physical Performance Battery [23], and Berg Balance Scale [24], and a lower prevalence of diabetes, less use of a walking aid, and faster gait speed.

Clinical measures

Sensory function

We used an abbreviated Semmes-Weinstein monofilament test (SWMT) to assess the threshold for light touch sensation on the dorsum of each great toe, which uses a buckling monofilament to impart a known force to the skin, following the protocol suggested by Perkins et al. [25]. The dorsum of the toe was used to avoid callouses on the plantar surface, which interfere with the sensory stimulus. The SWMT is a diagnostic tool to evaluate loss of protective sensation that often leads to ulcer formation. We employed two of the most-widely used monofilaments: the 5.07 monofilament (providing a standardized 10 g buckling force) and the 4.17 monofilament (providing a standardized force of 1.4 g). Failure to feel the 5.07 monofilament represents a loss of protective sensation [26, 27]. The 4.17 monofilament is used to determine whether the participant has normal light touch sensation. Inability to detect the 4.17 may indicate early neuropathy. The protocol began with the 4.17 monofilament, and if the participant could not feel it, he or she was tested with the 5.07 monofilament.

Monofilament testing was conducted while the participant was lying on an exam table, without shoes or socks. The procedure was demonstrated on the participant’s hand or arm before foot measurements were taken. Touch sensation was assessed at a single site on the dorsum of the great toe, 1 cm proximal to the nail bed (distal to the knuckle). The participant covered or closed their eyes throughout the test.

The test results were categorized into sensory loss groupings of intact, mild-to-moderate, and severe deficits, as defined in Table 1. The prevalence of each group is also shown. Changes in sensation over 5 years were grouped into 4 categories as shown in Table 2; including intact to intact, intact to mild-to-moderate impairment, mild-to-moderate impairment to severe, or severe to severe. Nine subjects who exhibited improved sensation were excluded from the longitudinal analysis.

Gait speed

Gait speed was assessed with a stopwatch as participants walked at their preferred pace over a four meter course. Timing started with a signal while subjects were standing still and ended when they traversed 4 m. To prevent terminal slowing, they were not told where the course ended. They were asked to walk at their comfortable speed as if taking a purposeful walk on the street, going to a store. They were allowed to use an assistive device if they used it at home or outdoors. The fastest time of two separate trials was used for analysis. The time to walk 4 m is a component of the Short Physical Performance Battery (SPPB), described below.
Table 1 The definition and prevalence of each category of somatosensory impairment

| Intact | Able to feel at least 3 monofilament touches out of 4 attempts for a 4.17 g monofilament in the left and right great toes | 292 (83%) |
| Mild-moderate impairment | Able to feel fewer than 3 monofilament touches out of 4 attempts for a 4.17 g monofilament in the left or right great toe and able to feel at least 3 monofilament touches out of 4 attempts for a 3.07 g monofilament in the left and right great toes | 22 (6%) |
| Severe impairment | Able to feel fewer than 3 monofilament touches out of 4 attempts for both 4.17 and 5.07 g monofilaments in the left or right great toe | 37 (11%) |

**Berg balance scale**
The Berg Balance Scale is a multi-component assessment of standing balance, consisting of 14 balance tasks with each task scored from 0 to 4, for a summed score of 0 to 56 [24]. The scale has been well-validated and shown to predict risk of falls in community dwelling elders [28]. Only baseline data were available.

**Physical performance**
The Short Physical Performance Battery (SPPB) was used to measure lower extremity mobility performance [23]. The SPPB includes measures of standing balance, 4-m usual-paced walking speed, and ability and time to rise from a chair 5 times. The validity of this scale has been demonstrated by showing a gradient of risk for admission to a nursing home and mortality along the full range of the scale from 0 to 12 [29, 30].

**Falls detection**
During a five-year follow-up period, a fall was defined as unintentionally coming to rest on the ground or other lower level, not as a result of a major intrinsic event (e.g., myocardial infarction, stroke, or seizure) or an overwhelming external hazard (e.g., hit by a vehicle) [18, 31]. Participants were instructed to complete and return monthly falls calendar postcards designed to be posted on a refrigerator. On the postcards, participants were to record an “F” for each fall on the day it occurred and an “N” on days when no falls occurred. If the postcard was not returned, a research assistant called the participant to determine whether a fall occurred during the preceding month and to remind them to complete and return future cards. This approach has been well-validated for use in epidemiological cohort studies and described in previous studies [32]. All subjects who reported falls were also called to determine the circumstances of the fall and clinical outcomes, including whether any injuries (e.g. fractures) and hospital visits occurred.

**Other variables**
Sociodemographic characteristics assessed at baseline in the home interview included age, sex, race (self-identified), and years of education. At each wave we used the validated Physical Activity Scale for the Elderly (PASE) to measure self-assessed physical activity in the previous week [33]. Participants were asked about physician-diagnosed major medical conditions.

Diabetes was defined using an algorithm based on self-reported diabetes, use of antidiabetic medications, and laboratory measures from the baseline clinic visit, including random glucose (> 200 mg/dL) and hemoglobin A1C (> 7%). Body mass index (BMI, calculated as weight in kilograms divided by height in meters squared) was calculated from measured height and weight. Comorbidity was measured using a count of relevant self-reported medical conditions, including: coronary heart disease, high blood pressure, ulcer or other stomach disease, kidney disease, liver disease, anemia, cancer, depression, osteoarthritis and degenerative arthritis, and rheumatoid arthritis [34]. Baseline subject characteristics also included the Center for Epidemiologic Studies Depression Scale – revised [35], Mini-Mental State Examination and Trail-making Test.

**Data analysis**
We summarized baseline characteristics of groups of participants with and without sensory loss using means and standard deviations or frequency distributions and compared groups of participants using t-tests for continuous variables, chi-square tests for categorical variables, and negative binomial regression for count variables.

Participants were grouped into 4 peripheral sensory loss categories based on their changes from baseline to 60 months: those with intact sensory function both at baseline and 60 months of follow-up (hereafter called consistently intact), those with intact sensory function at baseline and loss of function to at least mild impairment
by 60 months (progressing to impairment), those with at least mildly impaired sensory function at both baseline and 60 months (consistently impaired), and those with impaired sensory function at baseline but intact function at 60 months (improved) (Table 2). Those with mild-to-moderate and severe sensory loss were grouped together into the impaired category for the derivation of these sensory loss trajectories. Due to the small number of participants (9) in the improved category, this group was excluded from consideration in longitudinal analyses, leaving 342 participants for the analyses of change over time.

Analysis of the relationship between sensory loss categories and change in gait speed was done using repeated measures analysis of variance models. Gait speeds measured at baseline, 18 months, and 60 months were used as dependent variables, and tests of global mean differences across all time points as well as pairwise tests comparing mean gait speeds at each time point were conducted. The relationship to the falls rate outcome was analyzed using negative binomial regression, as falls rates exhibited a high degree of variance between participants through the course of study follow-up. Since falls were recorded as present or absent on each day of the 5-year follow-up period, there were no missing falls data.

Adjusted analyses for gait speed included age, sex, comorbidity count, use of a walking aid, baseline Berg balance score, and diabetes as covariates. Adjusted analyses for falls included age, sex, average weekly physical activity (PASE), baseline Berg balance score, diabetes, comorbidity count, and the SPPB score. For hypothesis testing, a two-sided type-I error probability of 0.05 was allowed. Models were estimated using Stata/MP version 13.1 (Statacorp, College Station, Tx).

Results
The characteristics of the full study cohort and those with (mild, moderate, or severe) and without (intact) baseline sensory impairment are shown in Table 3. At baseline, participants with any sensory impairment were four years older on average than their intact counterparts, were more likely to be male, and had diminished executive function (as indicated by 20s longer Trails B time on average). They also exhibited more multimorbidity and diminished physical functioning than their intact counterparts, reporting a greater prevalence of diabetes and peripheral neuropathy, and 0.3 greater mean comorbidity count on average. They were twice as likely to report use of a walking aid and exhibited diminished physical function, scoring an average of 1.4 points lower on the SPPB and 0.1 m/s slower gait speed, both differences well above established thresholds of clinical significance. Despite these differences, however, a history of one or more falls was similar in the two groups.

The relationship between changes in peripheral sensory function and change in gait speed over 18 and 60 months is illustrated in Fig. 1. After model adjustment, participants in the three groups had comparable mean baseline gait speed. The group of subjects with consistently intact sensory function over this time period had the smallest declines in gait speed over 60 months (-0.12 m/s; 95% CI: -0.15 to -0.08). This decline in gait speed was of similar magnitude to the decline observed in subjects whose sensory function progressed to impairment (-0.13 m/s; -0.16 to -0.10). Those whose sensory function was consistently impaired had a steeper decline in gait speed (-0.23 m/s; 95% CI: -0.28 to -0.18). The difference between the consistently impaired group and the others was statistically significant and consistent with a ‘substantially’ meaningful difference for gait speed metrics in older populations [36].

Declines in peripheral sensory function were also associated with fall risk. Table 4 shows the absolute and relative risk of falls by temporal pattern of impairment. Those whose sensory function progressed to impairment during follow-up had a greater risk of falls than those whose sensory function was consistently intact (adjusted risk ratio = 1.57 (95% confidence interval = 1.12 to 2.22). The group that remained consistently impaired over the 5 years had an elevated fall rate, but it was not statistically significantly different from that of the consistently intact group (adjusted risk ratio = 1.47, 95% confidence interval = 0.89 to 2.45).

Discussion
The results of this study demonstrate the course of tactile sensory loss in the feet of older community-dwelling people over 5 years, and relationships between peripheral sensory loss and the concurrent slowing of gait and development of falls. Over the 5-year course of the study, older adults with consistently impaired peripheral sensory function had a significantly greater decline in gait speed compared to those with consistently intact or progressive impairment over 5 years. Compared to those with consistently intact sensory function, older adults who developed sensory impairments had a greater risk of falls. These results indicate that the loss of peripheral sensory function is a significant contributor to slowing of gait and an increased risk of falls in a community-dwelling older adult population.

The absolute number (59) and percent (17%) of participants with sensory impairment were relatively low in the MOBILIZE Boston population, but higher than the 7% reported in other older populations [11, 13]. Most of our participants had intact sensation at baseline, but this enabled us to observe the development of impairments over time. Nearly half (49%) of those initially intact developed sensory impairment over 5 years of followup.
**Table 3** Baseline characteristics and descriptive statistics of the study sample (N = 351); mean (standard deviation) or count (percent) is shown

|                          | Full cohort (n = 351) | Baseline somatosensory function | Intact (n = 292) | Impaired (n = 59) |
|--------------------------|-----------------------|---------------------------------|------------------|------------------|
| Age, years               | 78 (5)                | 77 (5)                          | 81 (5)           |                  |
| Female                   | 230 (66%)             | 203 (70%)                       | 27 (46%)         |                  |
| White                    | 283 (81%)             | 230 (79%)                       | 53 (90%)         |                  |
| Education                |                       |                                 |                  |                  |
| Less than high school    | 19 (5%)               | 15 (5%)                         | 4 (7%)           |                  |
| High school              | 68 (19%)              | 53 (18%)                        | 15 (25%)         |                  |
| Some college/college     | 136 (39%)             | 119 (41%)                       | 17 (29%)         |                  |
| Graduate/professional    | 127 (36%)             | 104 (36%)                       | 23 (39%)         |                  |
| Body mass index, kg/m²   | 27.2 (4.9)            | 27.0 (4.9)                      | 27.9 (5.0)       |                  |
| Current smoker           | 11 (3%)               | 10 (3%)                         | 1 (2%)           |                  |
| Daily alcohol use (percent yes) | 34 (10%) | 28 (10%)                     | 6 (10%)          |                  |
| Comorbidity counta       | 2.4 (1.5)             | 2.4 (1.5)                       | 2.7 (1.5)        |                  |
| CES-D score              | 9.9 (10.2)            | 9.8 (10.0)                      | 10.3 (11.0)      |                  |
| Medication use           |                       |                                 |                  |                  |
| Antihypertensive medicine| 237 (68%)             | 194 (66%)                       | 43 (73%)         |                  |
| Antidepressants          | 35 (10%)              | 29 (10%)                        | 6 (10%)          |                  |
| Anti-seizure medications | 12 (3%)               | 10 (3%)                         | 2 (4%)           |                  |
| Statins                  | 160 (46%)             | 131 (45%)                       | 29 (49%)         |                  |
| Anxiolytics              | 42 (12%)              | 33 (11%)                        | 9 (15%)          |                  |
| Antihistamines           | 35 (10%)              | 29 (10%)                        | 6 (10%)          |                  |
| Opioids                  | 14 (4%)               | 10 (3%)                         | 4 (7%)           |                  |
| Nonsteroidal anti-inflammatory drugs | 65 (19%) | 51 (18%)                     | 14 (24%)         |                  |
| Analgesics/antipyretics  | 82 (24%)              | 71 (25%)                        | 11 (19%)         |                  |
| Cognitive function       |                       |                                 |                  |                  |
| Mini-Mental State Exam.  | 27.7 (2.3)            | 27.8 (2.3)                      | 27.5 (2.3)       |                  |
| Trail Making Test, seconds |                   |                                 |                  |                  |
| Part A                   | 52 (33)               | 51 (33)                         | 56 (33)          |                  |
| Part B                   | 123 (68)              | 120 (66)                        | 140 (72)         |                  |
| Part B less A            | 74 (55)               | 72 (55)                         | 84 (55)          |                  |
| Medical conditions (self-report) |               |                                 |                  |                  |
| Stroke                   | 32 (9%)               | 26 (9%)                         | 6 (10%)          |                  |
| Diabetes mellitus        | 47 (13%)              | 33 (11%)                        | 14 (24%)         |                  |
| Hyperlipidemia           | 179 (51%)             | 148 (51%)                       | 31 (53%)         |                  |
| Hypertension             | 259 (75%)             | 214 (74%)                       | 45 (78%)         |                  |
| Peripheral artery disease| 29 (8%)               | 22 (8%)                         | 7 (12%)          |                  |
| History of back pain or spinal stenosis | 138 (39%) | 116 (40%)                     | 22 (37%)         |                  |
| History of falls         | 129 (37%)             | 106 (36%)                       | 23 (40%)         |                  |
| Parkinson’s disease      | 6 (2%)                | 5 (2%)                          | 1 (2%)           |                  |
| Peripheral neuropathy    | 69 (21%)              | 35 (13%)                        | 34 (63%)         |                  |
| Cancer, excluding skin cancer | 81 (23%) | 63 (22%)                     | 18 (31%)         |                  |
This high incidence of sensory loss has not been fully appreciated in other studies nor in clinical practice. To our knowledge previous studies have not examined changes in peripheral sensory function over time nor their relationship to mobility outcomes. However, several cross-sectional and longitudinal studies have examined relationships between baseline abnormalities in foot sensation (including sense of vibration and touch), functional impairments, and fall risk. Most of these studies demonstrated that the loss of vibratory sensation was associated with increased fall risk [5, 37, 38]. For example, Patel et al. [37] found among older women with diabetes that fallers had higher vibratory sensory thresholds than non-fallers, but both fallers and non-fallers had similar touch sensation. Using cross-sectional data from the 2003–2004 National Health and Nutrition Examination Survey (NHANES), Wilson et al. [38] found no association between sensory loss in the feet, as measured using 10 g SWMT, and a subjective report of “difficulty with falls during the past 12 months,” assessed by questionnaire. However, after adjustments, participants who exhibited peripheral neuropathy had an increased risk of balance impairment compared to those without peripheral neuropathy. In a cross-sectional study exploring the effect of different sensory systems on postural stability, Lord and colleagues [5] found that poor tactile sensitivity to Semmes-Weinstein monofilaments at the lateral malleolus of the dominant ankle was associated with increased body sway. They also reported that peripheral sensation is the most important sensory system in the maintenance of static postural stability. Richardson et al. [39] found a cross-sectional relationship between peripheral neuropathy diagnosed by electromyography in a small referral population and a self-reported history of falls during the past year. Using longitudinal data from the Health, Aging and Body Composition study, Strotmeyer et al. [16] found that insensitivity to the 10 g Semmes Weinstein monofilament at the great toe was associated with lower quadriceps muscle strength. Thus, a number of neuromuscular deficits are associated with peripheral neuropathy and may lead to mobility impairments and falls.

Our longitudinal results are most consistent with those of Lord et al. [7] and Luukinen et al. [8] who found relationships between various modalities of sensory loss at the knees and recurrent falls in community-dwelling populations followed prospectively with fall questionnaires or diaries over 1 or 2 years, respectively. In Lord et al’s study, touch sensation did not

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Table 3 Baseline characteristics and descriptive statistics of the study sample (N = 351); mean (standard deviation) or count (percent) is shown (Continued)

| Physical function | Full cohort (n = 351) | Baseline somatosensory function |
|-------------------|----------------------|---------------------------------|
|                   |                      | Intact (n = 292) | Impaired (n = 59) |
| Uses walking aid  | 36 (10%)             | 24 (8%)           | 12 (20%)          |
| Physical Activity Scale for the Elderly | 111 (69) | 114 (70) | 97 (61) |
| Short Physical Performance Battery | 9.8 (2.2) | 10.0 (2.0) | 8.6 (2.8) |
| Gait speed at baseline, m/s | 1.00 (0.25) | 1.02 (0.24) | 0.92 (0.29) |
| Falls during first year of follow-up | 1.1 (1.7) | 1.1 (1.8) | 1.0 (1.4) |
| Berg Balance score | 51 (5) | 52 (5) | 48 (8) |

*Comorbidity count includes coronary heart disease, high blood pressure, diabetes, ulcer or other stomach disease, kidney disease, liver disease, anemia, cancer, depression, osteoarthritis and degenerative arthritis, rheumatoid arthritis, and other unlisted medical problem [34]
discriminate non-fallers from single or recurrent fallers. However, in another one-year prospective study of older adults living in an intermediate care facility, Lord and Clark [40] found that tactile sensitivity using a Semmes Weinstein Pressure Aesthesiometer did discriminate between fallers and non-fallers.

The subjects of our study who exhibited consistent sensory impairment at baseline and at follow-up were not more likely to fall than those who were consistently intact. The lack of a statistically significant difference between these groups could be due to the relatively small size of the consistently impaired group (N = 50) or adaptive behaviors learned by people with chronic sensory loss. Those with baseline sensory impairment were more likely to be using walking aids, which may have protected them from falls. There was also a trend towards lower physical activity as measured by PASE, suggesting that this group may have had less time at risk for falls.

The strength of the current study lies in its 5-year longitudinal design and rigorous detection of falls using state-of-the-art falls calendars and follow-up phone calls. There are several weaknesses to the current study. Participants did not receive a full neuropathic evaluation and the monofilament assessment of sensory function is only a quick screen that is specific, but not very sensitive to neuropathic sensory loss [41]. However, this simple, widely available, bed-side test of sensation on the dorsum of the great toe was sensitive to changes over time and predictive of a decrease in gait speed and increase in fall rate, even in the absence of more rigorous assessments of vibratory, proprioceptive, and motor nerve function. Although observed relationships could be confounded by underlying diseases such as diabetes, which affect sensation, gait and falls, the findings persisted after multivariate adjustments.

Our longitudinal results provide validation of the belief that peripheral sensory loss is a risk factor for mobility impairment. They also suggest that a decline in touch sensation at the great toe over time may be a more important predictor of slow gait speed and falls than a baseline sensory deficit. Therefore, simple, repeated clinical assessments of sensory function in the feet of older adults may be helpful in identifying and treating those at risk of falls to prevent their morbid consequences.

**Table 4** Risk of Falls According to temporal pattern in sensory function; point estimates and 95% confidence intervals shown

| Absolute Risk of Fall | Relative Risk of Fall |
|-----------------------|-----------------------|
|                       | Events (py) | Rate per py | Unadjusted | Adjusted |
| Consistently intact   | 384 (474)   | 0.81        | Referent    | Referent  |
| Progressing to impairment | 591 (454)   | 1.31        | 1.60 (1.16 to 2.22) | 1.57 (1.12 to 2.22) |
| Consistently impaired | 181 (161)   | 1.12        | 1.39 (0.89 to 2.18) | 1.48 (0.89 to 2.45) |

PY = patient year
*Estimates obtained using negative binomial regression
+Adjusted for age, sex, physical activity (PASE index), diabetes, comorbidity, Short Physical Performance Battery score

**Abbreviations**
BMI: Body mass index; MOBILIZE: Maintenance Of Balance, Independent Living, Intellect, and Zest in the Elderly; PASE: Physical Activity Scale for the Elderly; SPPB: Short Physical Performance Battery; SWMT: Semmes-Weinstein monofilament test

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**Availability of data and materials**
The datasets used and/or analysed during the current study are available from the corresponding author on reasonable request.

**Authors’ contributions**
Conception and design of the work: LL, BM and TT; Acquisition, analysis, interpretation of data for the work: LL, BM, DH, II, JZ and TT; Drafting the work and revising it for important intellectual content: LL, BM, JZ and TT. All authors read and approved the final manuscript.

**Ethics approval and consent to participate**
The MOBILIZE Boston Study was reviewed and approved by the Hebrew SeniorLife Institutional Review Board (IRB). Written informed consent was obtained from each participant. The study was conducted according to the principles of the Declaration of Helsinki.

**Consent for publication**
Not applicable.

**Competing interests**
The authors declare that they have no competing interests.

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