Predicting the Risk of Mobility Difficulty in Older Women With Screening Nomograms

The Women's Health and Aging Study II

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Background: A major obstacle to screening for early mobility disability (ie, mobility difficulty), a major public health concern, is the lack of a method that identifies those who are at high risk. The goal of this study was to develop easy-to-use clinical nomograms for estimation of the probability of incident mobility difficulty.

Methods: We conducted a population-based prospective study using data from 266 high physically and cognitively functioning older women, aged 70 to 80 years, who were free of mobility disability at the baseline evaluation of the Women's Health and Aging Study II. The outcome measure was incident mobility disability within 18 months, defined as self-reported difficulty walking 0.8 km, climbing 10 steps, or transferring from or into a car or bus. Logistic regression and receiver operating characteristic curve analyses were used for evaluation of the optimal combination of self-reported and performance-based mobility measures. Bootstrap sampling and estimation was used for validation.

Results: Predictive nomograms were developed based on a final model that included 3 simple-to-obtain measures of preclinical disability: self-report of modification in mobility tasks without having difficulty with them, one-leg stance balance, and time to walk 1 m at a usual pace. Final model accuracy (as estimated by the area under the receiver operating characteristic curve) was 73% (SE=0.04). Validation analysis confirmed the high accuracy of these nomograms.

Conclusions: An original tool was developed for assessment of the risk of mobility difficulty in older women that can be used to assist physicians and researchers in deciding which women to target for preventive interventions.

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OBILITY DISABILITY (ie, mobility difficulty) is a highly prevalent public health concern. Up to 50% of persons aged

65 years and older have disability in mobility-related tasks such as walking 0.4 km, climbing steps, transferring, or doing heavy housework.¹ Mobility is but one of several types of physical disability. Nevertheless, it is a major risk factor for difficulty and dependency in other domains of physical functioning,²⁻⁴ causing decreased quality of life in older adults⁴⁻⁶ and substantial social and health care needs.^{7,8} Consequently, prevention or postponement of mobility disability is a high priority.^{9,10}

Screening of older adults for risk of mobility difficulty is an important step toward prevention. Identification of individuals at highest risk could provide relevant information for targeting those who are, theoretically, most likely to benefit from preventive interventions. However, how to identify the subset of older individuals at highest risk is yet to be established. In the clinical setting, for example, physicians have to rely on their subjective impressions when assessing patients' risk of mobility difficulty. Effective methods for screening are needed.

During the past decade, extensive methodological and clinical research have laid the groundwork for the screening of older adults for risk of mobility difficulty. Important for the development of such a basis were (1) accumulation of evidence supporting the hypothesis of the existence of a preclinical stage of disability in the natural history of disablement, in which nondisabled older adults have "nonsymptomatic" critical decline in physical function that constitutes a major risk factor for progression to mobility difficulty¹¹⁻¹³ (such evidence suggests that preclinical mobility decrements in many cases precede overt mobility difficulty and are a marker of a high-risk group for whom interventions might potentially yield the greatest benefits in terms of decreasing the

PARTICIPANTS AND METHODS

STUDY POPULATION

The WHAS II is a population-based prospective study of 436 women aged 70 to 80 years with no or minimal physical disability living in the community. The WHAS II was designed to be a companion study to the WHAS I, a study of 1002 women representative of the one third most disabled women older than 65 years living in the community.1 The WHAS II sample was based on 3 replicate age-stratified random samples drawn from the Health Care Financing Administration Medicare files that listed all female beneficiaries who were 70 to 79 years old in 12 contiguous ZIP code areas in eastern Baltimore City and Baltimore County, Maryland, on March 1, 1994, October 1, 1994, and May 1, 1995. The methods of this study have been described previously.13 Briefly, a screening interview was used to assess scores on the Mini-Mental State Examination,²⁶ a widely used measure of cognition, and to determine difficulty in 4 domains of physical functioning²⁷: mobility; upper extremity movement; performance of tasks indicative of complex functioning, such as instrumental activities of daily living (IADLs)²⁸; and basic self-care tasks (Table 1). Eligibility for WHAS II consisted of difficulty in no more than one domain and a Mini-Mental State Examination score greater than 24. Of 1630 women screened, 880 were eligible for participation, and 436 agreed to participate in WHAS II. This apparent low initial response reflects the relatively low telephone recruitment rates²⁹ (telephone recruitment was used because of fiscal constraints) and the difficulty in getting older adults to participate in an intensive prospective study. Compared with WHAS II participants, those who were eligible but decided not to participate had less education and lower income and were more likely to rate their health status as fair or poor. This study considers the 266 participants who were free of mobility disability at baseline.

DATA COLLECTION

Study participants underwent a comprehensive baseline evaluation at the Johns Hopkins Functional Status Laboratory, Baltimore, Md, between August 1994 and February 1996, and then returned to the clinic for a similar follow-up assessment visit 18 months later. Trained interviewers conducted standardized face-to-face interviews and collected data on demographic characteristics, health behaviors, cognitive and functional status, and medical history. Two independent experts using standardized algorithms³⁰ subsequently validated the latter.

disability burden in older adults); (2) advances in the geriatric functional assessment field leading not only to identification of several risk factors for disability but, most important, to better characterization of functional impairments and their role in the mobility disablement process¹⁴⁻¹⁸; (3) the report of promising interventions that positively modify intermediate end points in the mobility disability pathway, such as lower extremity strength, gait velocity, and postural stability¹⁹⁻²⁵; and (4) the fact that women aged 75 years and older have the highest prevalence of chronic diseases and disability, indicating that this group might warrant special attention in terms of screening.^{5,7} The next challenge, then, is to translate Participants also underwent a thorough physical examination and had blood samples drawn. The procedures used were approved by the institutional review board of The Johns Hopkins Medical Institutions. Informed consent was obtained from all participants.

PHYSICAL FUNCTION ASSESSMENT

Details for obtaining the 3 measures for risk assessment using nomograms are provided in **Table 2**.

Self-reports of difficulty were obtained for 3 mobilityrelated and 24 other activities of daily life via questions such as, "For health or physical reasons, do you have difficulty climbing up 10 steps?" Self-reports of task modification in mobility tasks, a novel marker of preclinical disability, were obtained by asking individuals who reported no difficulty in mobility tasks whether they had modified the way they were performing them, by changing the method or reducing the frequency of task performance, because of underlying health problems. Recently, Fried et al¹³ demonstrated that this is a strong, independent predictor of developing incident mobility difficulty within 18 months that correlates well with performance-based measures of mobility functioning. Besides, it has good reliability (sameday test-retest weighted $\kappa = 0.74$).¹²

Objective and standardized performance-based measures were adapted from WHAS I protocols³¹ and included (1) time to walk 1 m at usual and rapid paces; (2) time to rise from a chair 5 times as rapidly as possible with arms crossed in front of chest; (3) one-leg stance balance,16,32 based on the length of time participants were able to maintain balance while standing on one leg up to a maximum of 30 seconds (using 3 categories: <10 seconds, 10 to <30 seconds, and 30 seconds); (4) the static balance measure, created by Guralnik et al,33 based on the participant's ability to maintain their feet in the side-by-side, semitandem, and tandem positions for 10 seconds; (5) maximal strength of the hip flexor muscles, assessed using a handheld dynamometer (2 trials for each leg were carried out, and the average strength of the strongest leg was used for analysis); (6) maximal grip strength of the dominant and nondominant hands using a handheld Jamar dynamometer (2 trials for each hand were carried out, and the average was used as the final measure for each hand); and (7) the Mini-Mental State Examination score. The intraclass correlation coefficient for the time to walk 1 m at a usual pace for measures obtained 1 week apart in WHAS I, which used the same protocol, was 0.90 (95% confidence interval, 0.89-0.91) and confirms the high reliability of this measure (data not published). Intraclass

such findings into new and clinically relevant approaches to screen for individuals with preclinical changes in mobility functioning who are at high risk for mobility difficulty.

In this study, we used data from the Women's Health and Aging Study II (WHAS II) to develop and validate an easy-to-use screening tool for estimation of the probability of incident mobility difficulty within 18 months in nondisabled community-dwelling women aged 70 to 80 years. The WHAS II was primarily designed to study the transition from preclinical to clinical disability, thus offering a unique opportunity for the development of a screening tool for early mobility disability. correlation coefficients for repeated chair stands time and grip and hip strength were also recently reported (0.76-0.89,³⁴ 0.92-0.95,³⁴ and 0.93,³⁵ respectively).

PRIMARY OUTCOME

Incident mobility difficulty within 18 months was the outcome of interest. For the purposes of these analyses, all participants at baseline were nondisabled, ie, they reported no difficulty in each of 3 tasks: walking 0.8 km, climbing 10 steps, and transferring from and into a car or bus. Those reporting difficulty in at least 1 of the 3 tasks at the 18month follow-up visit were considered to have incident mobility difficulty. The rationale for using these 3 tasks in our disability definition was based on (1) our clinical judgment that the broad concept of mobility functioning includes different components that have distinct biomechanical and energy demands that are best captured by 3 tasks instead of 1 and (2) previous work showing that these tasks are likely to be affected early in the mobility disablement process and constitute different and nonhierarchical entry points into this process (P.H.M.C., E.S.G., and L.P.F., unpublished data, 1994-1996).

STATISTICAL ANALYSIS

Step 1: Data Exploration

Descriptive statistics were used to characterize the study population and measures of physical function. Bivariate associations between incident mobility difficulty and potential predictors were examined using scatter plots, box plots, cross-tabulations, and logistic regression analyses, with and without adjustment for age. Threshold and nonlinear relationships were evaluated using quadratic and spline³⁶ terms during modeling (F and χ^2 statistics at α =.05 were used for evaluation of statistical significance).

Step 2: Multiple Logistic Regression

Modeling was carried out for selection of the most parsimonious combination of predictors of incident mobility difficulty 18 months later. The goal in using such an approach was to maximize data collection efficiency in the clinical setting. Only variables that achieved a significance level of P<.05 in step 1 were considered. A stepwise model was fit. Variables were entered into the model in an order that would reproduce the information collection process most likely to occur in the clinical setting. Likelihood

RESULTS

STUDY POPULATION

Study participants consisted of 266 communitydwelling women aged 70 to 80 years who reported no difficulty in 3 mobility tasks (**Table 3**). Almost 50% lived alone. Overall, there was significant variability in education, race, perceived health status, and number of chronic diseases. The most frequently diagnosed diseases were hypertension and osteoarthritis. Participants were cognitively intact. Task modification (ie, decrease in frequency or change in the method of performance of ratio tests were used to assess whether the addition of variables would provide supplemental information above and beyond that already obtained by a simpler model (P<.05 is necessary for statistical significance).

Step 3: Receiver Operating Characteristic Curve Analyses

Receiver operating characteristic curves and their respective areas under the curve (AUCs) were used for estimation of the overall accuracy of different models in predicting disability.³⁷ Differences in AUCs were then compared according to the method developed by Hanley and McNeil.³⁸

Step 4: Nomograms

Taking into account results from the receiver operating characteristic curve analyses and clinical feasibility, one model was identified as optimal for developing predictive nomograms. We then constructed fitted plots of the predicted probability of incident mobility difficulty, and their associated 95% confidence intervals, given the measures included in this optimal model. By connecting the points in these plots, we obtained the probability and confidence bands that define our nomograms.

Step 5: Validation Analysis

The prediction rule used for developing these nomograms was internally validated by taking 1000 bootstrap samples³⁹ from the original cohort and determining the median and 95% confidence intervals of 5 accuracy measures: sensitivity, specificity, positive and negative predictive values, and correct classification (ie, the percentage of individuals whose predicted and observed probabilities are the same). Briefly, fitted probabilities were computed according to the nomogram prediction rule and compared with the observed probabilities of incident disability. The latter assumed values of 1 or 0 depending on whether the individual developed or did not develop disability, respectively. For comparison, the fitted probabilities were dichotomized: they were defined as 0 if the fitted were lower than the probability cutoff point or 1 if higher. For example, if the computed probability was 0.40, her predicted probability was considered 0 for cutoff levels less than 0.40 and 1 for cutoff levels greater than 0.40. Bootstrap analyses were then conducted considering different cutoff points. Stata 6.0⁴⁰ and S-Plus 3.3⁴¹ were used for analyses.

mobility tasks) while having no difficulty with mobility tasks was reported by 26% of individuals. Objective performance-based measures of function were compatible with a high-functioning group.⁴²

DERIVATION OF NOMOGRAMS

After 18 months, 23.9% of study participants developed mobility difficulty. **Table 4** shows that incident mobility difficulty was predicted by self-report of task modification, time to walk 1 m at a usual or rapid pace, time to complete 5 chair stands, and one-leg stance balance. Variables that were not statistically significant predicTable 1. Domains of Physical Disability in Which Difficulty Performing Certain Tasks Was Used for Screening of Participants Into Women's Health and Aging Study II

Domain of Physical Function	Task	
Mobility	Walking 0.4 km, walking up 10 steps without resting, getting into and out of a bed or chair, doing heavy housework	
Upper extremity movement Complex functioning Self-care	Raising arms up over head, grasping or handling, lifting something as heavy as 4.5 kg Using the telephone, doing light housework, preparing meals, shopping for personal items Bathing or showering, dressing, eating, using the toilet	

tors included static balance summary score, hip and grip strength, and Mini-Mental State Examination score. Age-adjusted and unadjusted analyses yielded almost identical results.

Table 5 shows the results of forward stepwise logistic regression analyses to identify the most predictive, yet parsimonious, combination of measures that predicted mobility difficulty within 18 months. Because self-report of modification of task performance can be easily obtained through clinical interview and this measure strongly predicted incident mobility difficulty in step 1, it was included in all models. Walking and chair stand times were entered next. Time to walk 1 m at a usual pace and time to complete 5 chair stands added significant and nonredundant information to models containing only self-report of task modification (models 2 and 4; P=.006 and .003, respectively); time to walk 1 m at a rapid pace did not (model 3; P = .09). Time to complete 5 chair stands did not significantly improve fit over a model with task modification and time to walk 1 m at a usual pace (model 5; P = .11). On the other hand, addition of oneleg stance balance to models 2 and 4 resulted in significant improvement in fit (models 6 and 7; P=.002 for both). No interaction terms were statistically significant (data not shown). Based on these results, models 6 and 7 seemed to be the best final model candidates.

Figure 1 displays receiver operating characteristic curves for models 6 and 7 and for the simpler models 1 and 2 (the latter 2 were included for illustration of the gain in accuracy obtained with incorporation of additional measures). The AUCs were 73%, 72%, 70%, and 62%, respectively (SE=0.04 for all). The difference between AUCs of models 2 and 1 was 8% but achieved only marginal statistical significance (P=.08). On the other hand, models 6 and 7 represented a statistically significant gain in predictive accuracy over the simplest model 1 (11%; P=.02 and 10%; P=.03, respectively). The difference between the AUCs of models 6 and 7 was not significant (1%; P=.40).

ASSESSING THE RISK OF INCIDENT MOBILITY DISABILITY WITH NOMOGRAMS

Taking into account the AUC of each model and clinical feasibility issues (eg, asking an older adult to walk 1 m and timing it is easier to implement than the 5 chair stands test; besides, the latter also depends on chair specifica-

Table 2. Obtaining the 3 Measures for Risk Assessment Using Nomograms

Self-report of mobility task modification

- Presence or absence of mobility task modification among those who report no mobility difficulty can be obtained based on the answer to the following questions "For health or physical reasons, do you have any difficulty in (*task*)?", where tasks are (1) walk 0.8 km, (2) climb 10 steps, and (3) transfer from/into car/bus. Possible answers are "yes" or "no."
- 2. Mobility modifications are assessed in steps.
 - Step 1: "In the previous set of questions, I was interested in whether you have difficulty with a task. Now, I will be interested in whether you have changed the way or changed how often you do a task. Due to a health or physical condition, have you changed how frequently you (*task*)?", where tasks were the same as above. Possible answers are "no," "yes, do it less frequently," or "yes, don't do it anymore."
 - Step 2. "Due to a health or physical condition, have you changed (modified) the method that you used to (*task*)?", where changing the method meant doing tasks slower, using body in a different position or using a different method of moving, using different types of clothing, or having omitted some parts of the task, or having changed the home. Possible answers are "no" or "yes."
- 3. Self-report of mobility task modification should be coded as follows. Subjects who answer "no" to the 3 questions in all 3 tasks are considered as "no report of task modification." If they answer "no" to the first question (task difficulty), but "yes" either to the frequency or method changes questions, they are considered as having mobility task modification.

One-leg stance balance

Help participants into the appropriate position: standing on one leg, while keeping the other leg off the floor, and holding arms out to the side. When the subject is in position and ready, instruct to let go of the clinician's arm, and begin timing. The timing should be stopped when the subject moves her foot, grasps the clinician for support, or when 30 s elapses. One trial for each leg should be performed. The following one-leg balance categories should then be created based on the participant's best performance (ie, the longer time between the 2 trials): (1) category 1: able to hold balance <10 s; (2) category 3: able to hold balance for ≥ 10 s, but <30 s.

Time to walk 1 m at usual pace

Place a tape on the floor to serve as the starting line. Place 2 other tapes on the floor 1 and 4 m away from the initial tape. Ask the subject to stand with both feet touching the starting line. Then ask the subject to walk across a 4-m course at their usual speed, just as they were walking down the street to go to the store. Tell the subject to start walking after the prompt, "Ready, begin!" The clinician should walk beside the subject. Timing should be started as the participant begins walking and stopped when the participant's toe crosses the 1-m mark.

tions and is more burdensome), model 6 was chosen as the final model. Based on this model, we constructed 6 predictive nomograms (**Figure 2**), which were stratified by whether participants self-reported modification or no modification in mobility tasks because of health problems and by one-leg stance balance categories. Using these nomograms, the probability of developing mobility difficulty within 18 months can be readily obtained for nondisabled women aged 70 to 80 years. For example, if a woman reports that she has no difficulty with mobility tasks but that she has decreased the frequency she walks 0.8 km or biomechanically changed the way she climbs steps, and on examination she is able to stand on one leg for less than 10 seconds and she walks

Table 3. Demographic and Clinical Characteristics of the Study Population Serving as the Basis for the Derivation of Predictive Nomograms*

Baseline Category	Participants (N = 266)
Female sex	100
Age, mean ± SD (range), y	73.9 ± 2.8 (70-80)
Education, mean ± SD, y	12.7 ± 3.2
0-8	10.2
9-11	15.0
12	33.5
>12	41.4
Living alone	48.5
Race	
White	71.4
Black	27.1
Other	1.5
Perceived health status	
Excellent	7.5
Very good	13.9
Good	39.9
Fair	30.4
Poor	8.3
Chronic diseases	
Hypertension†	58.6
Osteoarthritis of the hip or knee‡	52.6
Osteoarthritis of the hand‡	34.2
Pulmonary diseases§	19.2
Osteoporosis	9.4
Coronary heart disease¶	8.6
Diabetes	5.3
Cancer	4.5
Stroke	2.3
Parkinson disease	0.4
Chronic diseases per person, mean ± SD, No.# Cognitive function	4.0 ± 1.2
MMSE score, median (5th, 95th percentiles)	29 (25, 30)

1 m at her usual pace in 2 seconds, her estimated probability of developing clinically significant mobility disability within 18 months is 56% (95% confidence interval, 40%-72%). Overall, risk is greater in individuals reporting task modification in mobility tasks decreases as individuals are able to maintain their one-leg balance for longer times, and increases with longer walking times.

VALIDATION ANALYSIS

Results from bootstrap analyses conducted for validation purposes are presented in **Table 6** and should be interpreted according to the following example. For the .30 cutoff point, the nomograms prediction rule have an estimated sensitivity of 44% (ie, the percentage of all the individuals who will develop mobility difficulty within 18 months have baseline fitted probabilities >.30), a specificity of 80% (ie, the percentage of those who will remain free of mobility difficulty have baseline fitted probabilities <.30), a positive predictive value of 40% (ie, the percentage of those whose baseline fitted probabilities were >.30 who will develop incident difficulty), and a negative predictive value of 83% (ie, the percentage of those whose baseline fitted probabilities were <.30 who will remain free of difficulty); overall correct classification will be achieved 72% of the time. Validity measures

Table 3. Demographic and Clinical Characteristics of the Study Population Serving as the Basis for the Derivation of Predictive Nomograms* (cont)

Baseline Category	Participants (N = 266)
Physical function	
Self-reported measures	
Difficulty in mobility tasks	0
Task modification in mobility tasks**	25.9
Performance-based measures	
Timed tests, median (5th, 95th percentiles), s	
Time to walk 1 m at a usual pace	1.4 (1.0, 2.0)
Time to walk 1 m at a rapid pace	1.0 (0.7, 1.4)
Time to complete 5 chair stands	12.5 (8.7, 17.8)
Time standing on one leg (up to 30 s)	10.1 (1.8, 30.0)
Balance	
One-leg stance balance	
Category 1 ($<$ 10 s)	49.2
Category 2 (10 to $<$ 30 s)	31.8
Category 3 (30 s)	19.0
Static balance ⁺⁺	
Score 1	3.4
Score 2	6.8
Score 3	20.4
Score 4	69.4
Strength, median (5th, 95th percentiles), kg	
Hip strength	16.6 (9.1, 29.3)
Grip strength—dominant hand	24.0 (17.3, 31.3)
Grip strength—nondominant hand	22.0 (15.3, 29.0)

*Data are given as percentages unless otherwise indicated. MMSE indicates Mini-Mental State Examination.

†Based on self-report of "diagnosis being told by a doctor." ‡Symptomatic or asymptomatic.

§Including those with chronic obstructive pulmonary disease and/or asthma.

||Including those with a diagnosis of osteoporosis and/or hip fracture. ¶Including those with angina and/or myocardial infarction (asymptomatic or symptomatic).

#Of the 10 adjudicated diseases listed.

** Participants who reported no mobility difficulty but had modified the way they performed mobility tasks by changing the method or reducing the frequency, because of underlying impairments.

[†]†Created according to Guraľnik et al³⁴: score 1 (the worst), able to stand in side-by-side position for 10 seconds but unable to stand in semitandem position for < 10 seconds; score 2, able to stay in semitandem position for 10 seconds but in full tandem position for <3 seconds; score 3, able to stay in semitandem position for 10 seconds and in full tandem position for 3 to < 10 seconds; and score 4 (the best), able to stand in full tandem position for 10 seconds.

are also shown for other cutoff points. As shown in Table 6, correct classification of individuals regarding future difficulty status can be expected 72% to 79% of the time for cutoff points of .30 or higher. In our study, 25% of older women who were free of mobility difficulty at baseline had fitted risks greater than .30.

COMMENT

We presented nomograms for straightforward estimation of the probability of developing mobility difficulty within 18 months for nondisabled, communitydwelling women aged 70 to 80 years. To estimate such risk, 3 measures need to be obtained and "plugged" into the nomograms: (1) self-report of whether the individual has changed the method or frequency of performing mobility tasks, while having no difficulty with them;

Table 4. Crude and Age-Adjusted Odds Ratios (ORs) With 95% Confidence Intervals (CIs) for Incident Mobility Difficulty Within 18 Months by Individual Physical Function Measures at Baseline

	Douticinanto No *	Unadjusted		Age Adjusted	
Measure of Physical Function	(N = 266)	OR (95% CI)	Р	OR (95% CI)	Р
Self-report of task modification (yes vs no)	266	3.16 (1.71-5.74)	.001	3.14 (1.71-5.74)	<.001
Time to walk 1 m at a usual pace (1 s)	256	4.47 (1.91-10.48)	.001	4.39 (1.86-10.37)	.001
Time to walk 1 m at a rapid pace (1 s)	263	4.41 (1.31-14.8)	.02	4.25 (1.25-14.50)	.02
Time to complete 5 chair stands (5 s)	258	2.26 (1.37-3.72)	.001	2.28 (1.38-3.77)	.001
One-leg stance balance†	258	· · · · · ·		()	
Category 1 (<10 s)		1.00		1.00	
Category 2 (10 to $<$ 30 s)		0.51 (0.26-0.98)	.04	0.51 (0.26-0.99)	.05
Category 3 (30 s)		0.09 (0.02-0.39)	.001	0.09 (0.02-0.39)	.001
Static balance [±]	265	· · · · · ·		()	
Score 1		1.00		1.00	
Score 2		1.27 (0.24-6.82)	.78	1.32 (0.24-7.11)	.75
Score 3		0.51 (0.11-2.38)	.39	0.52 (0.11-2.43)	.41
Score 4		0.57 (0.14-2.39)	.45	0.60 (0.14-2.53)	.49
Hip strength§ (1 kg)	259	1.00 (0.96-1.05)	.95	1.00 (0.96-1.05)	.93
Grip strength (1 kg)	261	0.98 (0.92-1.05)	.55	0.98 (0.92-1.05)	.65

*Number of individuals with measures available for each test. Data are not available for 0 to 10 people because of missing data or exclusion of outliers. †One trial on each leg was performed. Analyses are shown for the best performance, ie, the longer time.

‡For explanation, see the last footnote to Table 3.

§Two trials were performed on each leg; average strength for the stronger leg is used. Analyses were also carried out using the average strength for the weaker leg and the average between them, but results were virtually identical (data not shown).

||Presented is grip strength of the dominant hand. Analyses were also carried out for the nondominant hand and averaged, but results were virtually identical (data not shown).

Table 5. Multivariate Logistic Regression Models Used to Identify the Most Parsimonious Combination of Predictors of Developing Mobility Difficulty Within 18 Months

Model	Predictors Odds Radio (SE)		Likelihood Ratio Test*	
1	Self-report of task modification (yes vs no)	3.16 (0.97)	Not applicable	
2	Self-report of task modification (yes vs no) Walking time for 1 m at a usual pace (1 s)	2.83 (0.92) 3.36 (1.50)	Models 2 and 1: 7.49 (1 <i>df</i>); <i>P</i> = .006	
3	Self-report of task modification (yes vs no) Walking time for 1 m at a rapid pace (1 s)	2.99 (0.95) 3.02 (1.90)	Models 3 and 1: 3.84 (1 <i>df</i>); <i>P</i> = .08	
4	Self-report of task modification (yes vs no) Time to complete 5 chair stands (5 s)	3.13 (1.00) 2.17 (0.58)	Models 4 and 1: 8.67 (1 <i>df</i>); <i>P</i> = .003	
5	Self-report of task modification (yes vs no) Walking time for 1 m at a rapid pace (1 s) Time to complete 5 chair stands (5 s)	3.06 (1.00) 2.41 (1.20) 1.60 (0.47)	Models 5 and 2: 2.56 (1 <i>df</i>); <i>P</i> = .11	
6	Self-report of task modification (yes vs no) Walking time for 1 m at a usual pace (1 s) One-leg balance category 2 (10 to <30 s)† One-leg balance category 3 (30 s)†	2.37 (0.81) 2.23 (1.10) 0.60 (0.21) 0.13 (0.09)	Models 6 and 2: 12.4 (2 <i>df</i>); <i>P</i> = .002	
7	Self-report of task modification (yes vs no) Time to complete 5 chair stands (5 s) One-leg balance category 2 (10 to <30 s)† One-leg balance category 3 (30 s)†	2.55 (0.86) 1.89 (0.55) 0.59 (0.21) 0.12 (0.09)	Models 7 and 4: 12.9 (2 <i>df</i>); <i>P</i> = .002	

*Testing statistical significance of fit improvement resulting from the addition of a new variable(s). +Reference category is category 1 (<10 s).

(2) one-leg stance balance categories; and (3) time to walk 1 m at a usual pace. These 3 measures can be easily obtained in clinical settings and are safe, inexpensive, and not time-consuming (Table 2). From the functional assessment point of view, these measures are markers of preclinical disability¹³ and identify functional decline before it might be symptomatically apparent. The development of these nomograms provides an original screening tool for risk assessment of early and clinically relevant mobility disability in older women.

Previously, Guralnik et al¹⁴ used mobility-related performance-based measures to predict self-reported mobility difficulty. Specifically, that study tested how well a predefined combination of 3 performance-based measures predicted subsequent mobility dependency, a disability outcome measure shown to be hierarchically distinct from mobility difficulty.⁴³ The present study builds on that work by aiming to determine the most clinically useful combination of self-report and performancebased measures of mobility function for screening for risk



Figure 1. Receiver operating characteristic curves for assessing accuracy of elected logistic regression models in predicting incident mobility difficulty 18 months later. Model 6 (self-report of mobility task modification+time to walk 1 m at a usual pace+one-leg balance) and model 7 (self-report of task modification+time to complete 5 chair stands+one-leg balance) were the most effective and parsimonious combinations of predictors of incident mobility difficulty in step 2. These were compared with simpler model 2 (self-report of task modification+time to walk 1 m at a usual pace) and model 1 (self-report of task modification alone) to allow assessment of gain in accuracy resulting from including additional measures in the model. Predictive accuracies (defined in terms of the area under the curve) of models 6, 7, 2, and 1 were 73%, 72%, 70%, and 62%, respectively (SE=0.4 for all).

of mobility difficulty, an earlier stage of the disablement process. Regarding other screening tools, Moore and Siu⁴⁴ recently proposed the timed "Up & Go" test (the time for the individual to rise from an armchair, walk 3 m, turn, walk back, and sit down again). Although simple and appealing, it has yet to have its prospective screening accuracy established.

In addition to the major results of this study, other findings should be noticed. The static balance component of the tool developed by Guralnik et al¹⁴ was not a statistically significant predictor of mobility difficulty in our study, but the one-leg stance balance measure was. This suggests that the latter might be more sensitive and appropriate for the screening of higher-functioning populations. Also, time to complete 5 chair stands did not provide statistically significant information beyond that provided by 1-m walking time regarding short-term prediction of mobility difficulty. Finally, hip strength was not a significant predictor of incident mobility difficulty. This was likely because of a ceiling effect (ie, these high-functioning women's strength was beyond a threshold above which increases in strength do not add further protective effect).35

The decision to consider only measures of mobility functioning for inclusion in the prediction rule was based on 2 points: first, the conception that they represent not only the effects of chronic disease but also disease status– related deconditioning (ie, lower level of fitness) on functioning decline and should be among the most proximal measures to the outcome measure of this study, mobility difficulty, and second, an a priori expectation that use of only a few selected measures of mobility would allow reasonably accurate risk prediction, therefore obviating the need for additional measures from other domains. Indeed, increases in the time needed for obtaining such additional measures in the clinical setting and in the complexity for communicating risk would certainly represent



Figure 2. Nomograms to predict the probability of onset of mobility difficulty within 18 months in nondisabled women aged 70 to 80 years. Probability of incident mobility difficulty can be estimated directly from the nomograms using 3 simple measures (see Table 2): (1) self-report of mobility task modification (ie, asking whether, because of underlying impairments, individuals had modified the way they performed mobility tasks without having difficulty with them, by changing the method or the frequency of task performance), (2) one-leg stance balance (ie, the length of time the individual is able to maintain her balance while standing on one leg: <10 seconds, 10 to <30 seconds, or 30 seconds), and (3) time to walk 1 m at a usual pace. Left, Nomograms for individuals who report no mobility task modification. The rows are stratified by one-leg standing balance categories.

a threat to the clinical usefulness of our nomograms. Nonetheless, we conducted a post hoc analysis to evaluate the impact of 6 established risk factors for mobility difficulty not considered as candidates for our prediction rule on model fit and predictive accuracy: (1) definite diagnosis of angina or myocardial infarction, (2) definite diagnosis of knee osteoarthritis (symptomatic or asymptomatic), (3) presence of depression (\geq 11 points on the Geriatric Depression Scale),⁴⁵ (4) body mass index (<18.5 and >18.5 <25 and >25, and <30 and >30 kg/m²), (5) history of falls within the past 12 months, and (6) self-reported fear of falling within the past 12 months. When added to the prediction rule one at a time, none of these measures improved the predictive ability of our predictive rule in a statistically significant way (data

Probability Cutoff Point		Measures of Accuracy, %			
	Sensitivity	Specificity	Positive Predictive Value	Negative Predictive Value	Correctly Classified
≥.10	96 (90-100)	25 (8-31)	28 (21-34)	96 (89-100)	41 (35-48)
≥.15	95 (88-100)	37 (30-44)	31 (24-38)	96 (90-100)	50 (44-57
≥.20	79 (67-88)	55 (48-62)	34 (26-42)	90 (84-95)	60 (55-66
≥.25	61 (49-74)	72 (65-77)	39 (29-49)	86 (81-91)	69 (64-75
≥.30	44 (31-57)	80 (74-85)	40 (28-52)	83 (77-88)	72 (67-77
≥.35	39 (27-52)	83 (77-88)	41 (28-54)	82 (77-87)	73 (68-79
≥.40	30 (19-42)	91 (86-94)	49 (32-64)	81 (76-86)	77 (71-82
≥.45	21 (10-32)	93 (88-96)	46 (26-65)	80 (74-84)	76 (71-81
≥.50	16 (7-26)	97 (94-99)	60 (33-85)	79 (74-84)	78 (73-83)
≥.55	11 (4-19)	99 (97-100)	75 (40-100)	79 (74-84)	79 (74-84)

*Data are given as mean (bias corrected 95% confidence interval).

not shown). These findings can be explained in part by the relatively low variability in disease severity in highfunctioning populations.

The strengths of these nomograms include (1) prediction of incident mobility difficulty that is as accurate as current National Cholesterol Education Program guidelines and the single ratio of total plasma cholesterol level to high-density lipoprotein cholesterol level for prediction of coronary heart disease mortality (AUC=0.73, 0.74, and 0.72, respectively)⁴⁶ and as accurate as Papanicolaou smears for prediction of cervical cancer (AUC=70%)⁴⁷; (2) risk characterization as a probability, which is easier to communicate to patients than an odds ratio; and (3) provision of a range for the estimated probability.

Four important limitations of this study must be pointed out. First, initial nonresponse of individuals chosen for inclusion in this population-based cohort is likely to have limited the external validity of our results. Whether they can be generalized to other age-sex groups, which have overall lower risk of mobility disability than the population targeted here, or to less educated populations remains to be determined. Second, nonresponse is likely to have biased our results toward underestimation of the true risk of mobility difficulty onset, especially among those who reported mobility task modification or were classified in the worst categories of one-leg stance balance, considering that those who were eligible but decided not to participate had lower levels of functioning than did study participants. Third, validation of the predictive rule used for developing these nomograms, which showed a relatively high degree of predictive validity for the proposed screening tool, was internal rather than external. The latter is an inherently stronger approach, but given the nonexistence of another data set containing all the measures included in our prediction rule, and the relatively small sample size of this study, which could have led to significant loss of power and generation of unstable and nonprecise estimates had a sample-splitting technique been used, validation of the prediction rule in an independent population was not performed at this time. Finally, the confidence intervals presented in our nomograms do not incorporate day-to-day variability of the measures used in the prediction rule, such as 1-m walking time. Nonetheless, there is evidence in the literature that the size of these short-term fluctuations are relatively small, and individuals' locations in the population distribution of performance tend to be preserved for at least 6 months,⁴⁸ thus not constituting a threat for risk stratification purposes, which was the broad goal of this study.

In conclusion, this article offers a new approach to screening high-functioning older women for subsequent mobility difficulty. There are 2 contexts in which these nomograms have potential for application. First, they could be used for clinical screening and decision making. Evidence of a low disability risk can be used to reassure patients of their high functioning status, whereas further clinical evaluation might be sought for patients at high risk. Providing patients with knowledge about their short-term difficulty risk might encourage them to engage in the primary prevention activities currently recommended.⁴⁹⁻⁵² Second, by permitting identification of a preclinically disabled, highrisk group of older adults, these nomograms could contribute to the delineating effective preventive interventions for forestalling the cascade of functional decline that leads to the onset of early mobility disability.

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