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1 Different outcome measures and domains of functioning: 18

2 months follow-up of persons with dizziness

3 Abstract

4	Aims: To explore changes in different outcome measures in a follow-up of persons with
5	dizziness, and to investigate if these changes indicate different domains of functioning.
6	Methodology: Sixty-eight persons with dizziness, mean age of 47 years, were included
7	in an 18-months follow-up. Outcome measures used: the Vertigo Symptom Scale,
8	Patient Specific Functional Scale, Disability Scale, Dizziness Handicap Inventory, and
9	tests of Dynamic Visual Acuity, Single Leg Stance, and Walking Speeds.
10	Major findings: We found significant improvements in impairments indicated by
11	outcome measures of dizziness and visual acuity, and in activity and participation
12	indicated by outcome measures of standing balance, patient specific activities,
13	disability, and quality of life. Similar patterns of change were also found in subgroup
14	analyses, except in gender. Correlations between change-scores ranged $r = 0 - 0.6$.
15	Significant correlations were found between change scores indicating body function and
16	activity/participation ($0.3 \le r \ge 5$). We found no correlations between self-report
17	measures and tests.
18	Conclusion: Comprehensive use of outcome measures addressing body function,
19	activity and participation appear to provide information of changes in different domains
20	of functioning. To enable broad and meaningful follow-up of patients with long lasting
21	dizziness, generic and condition specific measures, self-reports and tests in different
22	domains of functioning could be further explored.

Key words: Vertigo Symptom Scale, Patient Specific Functional Scale, Disability
 Scale, Dizziness Handicap Inventory, Dynamic visual acuity

1 Background

2 The prevalence of dizziness in general populations mostly range between 16 and 30%, 3 dependent on definitions and methods used (1-4). Dizziness is associated with other 4 symptoms, conditions, diseases and use of medicines (3-5), and the prevalence increases 5 with the number of other symptoms and conditions, as well as number of medicines 6 used (3, 6). Dizziness often leads to impaired functional ability and quality of life (1, 7), 7 and accounts for a considerable use of health services (1). Occupational difficulties are 8 experienced by many (7, 8). Dizziness as such is a rare diagnosis of certified absence 9 from work, while absence of eight weeks or more due to this symptom, is found to 10 result in disability pension in about 25% after 5 years (9). 11 12 Studies indicate that about 20% to 70% of patients have persisting dizziness at 12 13 months follow-up or more (6, 8, 10, 11). Different patient groups, settings, length of 14 observations, and follow-up measures could explain the broad range. Predictors of 15 persistent dizziness are higher age; female gender; the origin of dizziness being 16 vestibular neuronitis and vestibular migraine; inactivity and avoidance of situations that 17 provoke dizziness; physical and mental co-morbidity; low self-rated health, and long 18 duration of symptoms at the time of inclusion (8, 10-14). 19 20 The World Health Organization (WHO) presented the International Classification of

Functioning, Disability, and Health (ICF) in 2001 (15) The theoretical model is neutral when it comes to causal relations between different domains of functioning. The health domains of the ICF comprise body structure and function (versus impairments), activity (versus limitations) and participation (versus restriction). Personal and environmental factors may have positive or negative effect on the health domains. WHO promoted the

1	use of outcome measures that address the different health domains and the use of
2	subjective and objective measures (15). Associations between scores in measures of
3	body function and structure, activity and participation are generally found to vary
4	between correlations of 0.2 and 0.5 (16). These findings indicate that result from
5	measures in one domain of functioning provides little information about other domains
6	of functioning (16). Based on this, one might assume that patients with persisting
7	symptoms and impairments still might improve in activity and participation.
8	
9	In 2000, experts (17) in patients with dizziness suggested relevant assessments and
10	outcome measures to these patients. In addition to insight in the history of dizziness and
11	personal and environmental factors, clinical tests and self-reports comprised symptoms
12	of dizziness, oculomotor and vestibular-ocular symptoms, proprioception,
13	musculoskeletal status, balance and ambulation, disability, health status, and quality of
14	life (17). The suggestions appeared to address body function/impairments, activity/
15	limitations, and participation/restrictions, as well as personal and environmental factors,
16	in line with the theoretical framework of ICF (15).
17	
18	In follow-ups \geq 12 months in patients with dizziness, only a few studies report changes
19	in different domains of functioning (8, 18, 19). Likewise, reports from follow-ups ≥ 12
20	months after vestibular rehabilitation comprise either self-reports or tests. Self-reported
21	results include symptoms of dizziness and disability (20, 21), self-perceived handicap
22	(22), and quality of life (23). Reports from tests include vestibular testing and/or tests of
23	balance/gait (24-26). Application of outcome measures that address different domains
24	of functioning as body structure and function, activity and participation, might provide
25	broader indications of long term changes of functioning in persons with dizziness.

1

2 The aim of this study is to explore changes in different outcome measures in a long term 3 follow-up of persons with dizziness, and to investigate if these changes indicate 4 different domains of functioning. 5 6 Methods 7 Study sample 8 Persons from the region of Oslo and Akershus were recruited between September 2003 9 and December 2004 from general practice, medical specialist, and the National 10 Insurance Administration (NIA). Inclusion criteria were patients with dizziness, age 11 range 20-65 years, and ability to read and understand Norwegian. Exclusion criteria 12 were individuals with recent cardio-vascular disease, neurological or other severe 13 system diseases, and/or not being able to go through physical tests. Of the 96 included, 14 63 were recruited from NIA, 24 from medical specialist, and 6 from general 15 practitioners. Three persons did not reply to the question of recruitment. Twenty-eight 16 persons were lost to follow-up at 18 months, giving a final sample of 68 participants. 17 Flowchart of participants and reasons for withdrawals is shown in Figure 1. Differences 18 between participants and withdrawals at initial testing did not reach statistical 19 significance as concerns characteristics, and average scores in questionnaires and tests. 20 21 All participants confirmed self-perceived dizziness and/or instability in the course of the 22 last 14 days according to Numeric Rating Scales. Description of participants is shown in 23 Table 1. About 2/3 were women, and the average age was 46.7 years. Time since onset

of symptoms had a median of about 1 ³/₄ years, and all participants had dizziness that

- had lasted \geq 4 months. Twenty-one persons (31%) were on varying percentage of sick
- 26 leave at inclusion. Sick-leave/social security benefit due to dizziness in the total group,

1	had a median duration of six months (Table 1). The origin of dizziness was gathered
2	from written medical information and self-reports, both presented by participants at
3	initial evaluation. The information on origins of dizziness was categorized as vestibular,
4	n = 44 (65%), non-vestibular, $n = 15$ (22%), and unknown, $n = 9$ (13%) (Table 2).
5	
6 7	<i>Procedure</i> The participants received self-administered questionnaires to be returned by mail prior
8	to an appointment for interview and testing. Follow-ups were administered at 6, and 18
9	months, using the same procedure. Here, only results from 18 months follow-up are
10	presented. The participants were informed about follow-up assessments, given
11	information about results from outcome measures, and suggestions concerning physical
12	activities. The same physiotherapist (the first author) interviewed and tested all
13	participants.
14	
15	Twenty-nine persons (43%) followed a 10 week group intervention program at the
16	
	Outpatient Department at the Physiotherapy Program, Oslo University College. The
17	Outpatient Department at the Physiotherapy Program, Oslo University College. The second author carried out the group intervention. The other 39 participants followed
17	second author carried out the group intervention. The other 39 participants followed
17 18	second author carried out the group intervention. The other 39 participants followed treatment as usual, which included persons who continued ongoing treatment elsewhere
17 18 19	second author carried out the group intervention. The other 39 participants followed treatment as usual, which included persons who continued ongoing treatment elsewhere uncontrolled by us, or got no specific treatment. The effect of intervention was not the
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 17 18 19 20 21 22 	second author carried out the group intervention. The other 39 participants followed treatment as usual, which included persons who continued ongoing treatment elsewhere uncontrolled by us, or got no specific treatment. The effect of intervention was not the focus of the present study.

according to the ICF, had not been established. The following generic and condition
 specific measures were applied:

3

Numeric Rating Scale (0-10 points) (27) on perceived symptoms of dizziness and
instability were used only at inclusion, where 0 indicated no dizziness/instability and 10
indicated the worse possible dizziness/instability perceived in the course of the last 14
days.

8

22

9 The Vertigo Symptom Scale – Short Form is a self-report questionnaire assessing 10 perceived severity of symptoms during the last month (28). It includes 15 items in two 11 subscales named "vertigo/balance" (8 items) and "autonomic/ anxiety" (7 items) (28). 12 Five ordinal response categories range from "never" (score 0) to "very often (most 13 days)" (score 4), with a total score ranging from 0 to 60 points (28). The Norwegian 14 version has satisfactory measurement properties (29). The smallest detectable change 15 (SDC) at group and individual level are ≥ 1.47 points (SDC_{group}), and ≥ 7.8 points 16 (SDC_{ind}) (29). Cut-off values of "severe dizziness" is defined as ≥ 12 points (30). The 17 Vertigo Symptom Scale was assumed to indicate body function/impairment according 18 to the ICF chapters Mental functions (b1), Sensory functions (b2), and 19 Neuromusculoskeletal and movement related functions (b7) (31). 20 21 The Patient Specific Functional Scale (32) is a generic scale of subjective functional

23 difficulties with, or is unable to perform because of her/his condition/problem, and rates

limitations. The patient is asked to list up to 5 important activities that she/he has

the degree of problems with each activity on an 11-point scale: 0 (cannot perform) to 10

25 (can perform as before injury/ disease/ illness). At follow-up, the patient is asked to

1 score the current level of difficulty associated with each of the activities (32). An 2 average score is estimated from the three first activities. Acceptable validity, test-retest 3 reliability and responsiveness to change have been reported in patients with different 4 dysfunction, but not in persons with dizziness (16). The smallest detectable change is 2 5 points in the average score and 3 points in a singular activity (90% CI) (16, 32). The condition was defined as "dizziness" in the present study, and the person was informed 6 about his/her initial score in each activity. The Patient Specific Functional Scale was 7 8 assumed to enable the participants to report components of activity/limitations and 9 participation/restrictions according to the ICF chapter d (31). 10 11 The Disability Scale assesses functional ability in connection with dizziness (33), with 12 one question: "Please mark the one statement that best describes your present situation".

Six ordinal response categories range 'No disability; negligible symptoms' (score 0) to 'Long-term severe disability; unable to work for over 1 year or established permanent disability with compensation payment' (score 5) (33). Reliability of the scale has been demonstrated in patients with peripheral vestibular disorders, and a change of only one category indicates a significant change in disability (34). The Disability Scale was assumed to indicate components of participation/restrictions according to ICF categories in Domestic life (d6) and Major life areas (d8) (31).

20

The condition specific *Dizziness Handicap Inventory* assesses the impact of dizziness on physical, functional and emotional domains, interpreted as indications of quality of life and functional ability (35). A total score (0-100) is obtained by summing ordinal scale responses to 25 questions (Yes = 4, Sometimes = 2, No = 0) (35). The Norwegian version has satisfactory measurement properties of the total scale (36). Cut-off point for

1	'disability' versus 'no disability' is 29 points. The smallest detectable change (SDC) at				
2	group and individual level are \geq 4 points (SDC _{group}), and \geq 20 points (SDC _{ind}) (36).				
3	Minimally important change (MIC) in the inventory is found to be ≥ 11 points (36), and				
4	indicates the ability of an instrument to detect clinically important change, a real change				
5	in the concept being measured (37). The Dizziness Handicap Inventory was assumed to				
6	indicate components in body function/impairment, as well as activity/limitations and				
7	participation/restrictions according to the ICF chapters in Mental functions (b1),				
8	Mobility (d4), Domestic life (d6), Major life areas (d8), Community, social and civic				
9	life (d9) (31).				
10					
10					
10 11	The Dynamic Visual Acuity Test measures the degradation of visual acuity that occurs				
	The <i>Dynamic Visual Acuity Test</i> measures the degradation of visual acuity that occurs with head movement, indicating the ability to see clearly during head movement (38). In				
11					
11 12	with head movement, indicating the ability to see clearly during head movement (38). In				
11 12 13	with head movement, indicating the ability to see clearly during head movement (38). In patients with uncompensated, unilateral vestibular loss, dynamic visual acuity is known				
11 12 13 14	with head movement, indicating the ability to see clearly during head movement (38). In patients with uncompensated, unilateral vestibular loss, dynamic visual acuity is known to degrade by three or four lines (39). An average improvement of more than 2 lines is				
 11 12 13 14 15 	with head movement, indicating the ability to see clearly during head movement (38). In patients with uncompensated, unilateral vestibular loss, dynamic visual acuity is known to degrade by three or four lines (39). An average improvement of more than 2 lines is reported following vestibular rehabilitation (40). We applied a clinical test, since the				
 11 12 13 14 15 16 	with head movement, indicating the ability to see clearly during head movement (38). In patients with uncompensated, unilateral vestibular loss, dynamic visual acuity is known to degrade by three or four lines (39). An average improvement of more than 2 lines is reported following vestibular rehabilitation (40). We applied a clinical test, since the more reliable computerized equipment was not accessible at our clinic. The number of				

 $20 \qquad \text{the ICF category b215 (31).}$

21

Single Leg Stance is a measure of standing balance (38). The average of two trials in each test was used, unless the participant reached 30 seconds in the first trial. In this study group the smallest detectable changes at group and individual levels were 1.45 seconds (SDC_{group}) and 12.53 seconds (SDC_{ind}) in the right leg, and 1.65 seconds

(SDC_{group}) and 14.04 (SDC_{ind}) in the left leg. The Single Leg Stance-eyes closed was
 assumed to indicate activity/limitation according to the ICF category Mobility d415
 (31).

4

5 Walking speed is commonly used as a measure of functional balance (16, 38). Preferred 6 walking speed has acceptable reliability in different patient populations (16), as well as 7 in patients with peripheral vestibular disorders (34). Reference values are provided in 8 healthy adults (41, 42). Fast walking speed is known to be a problem in patients with 9 dizziness. Tests of preferred and fast walking speeds were applied, the average of two 10 trials in each test was calculated to estimate average walking distance meters/second 11 (m/s). In this study group the smallest detectable change in preferred walking speed at 12 group and individual level were was 0.02 m/s (SDC_{group}) and 0.2 m/s (SDC_{ind}). This was 13 in line with previous reports (34). The smallest detectable change in fast walking speed 14 at group and individual level were 0.03 m/s (SDC_{group}) and 0.3 m/s (SDC_{ind}). Walking 15 speeds were assumed to indicate activity/limitation according to the ICF category 16 Mobility d450 (31). 17

18 Ethics

The study was recommended by the Norwegian Regional Committee for Medical
Research Ethics and the Norwegian Social Science Data Services. The study was
performed in accordance with the Helsinki Declaration. Written informed consent was
obtained from all participants.

23

24 Statistical analyses25 Questionnaires with

25 Questionnaires with missing values exceeding 30% were discarded, while missing values \leq 30% of the

26 items in a participant's questionnaire were imputed by the mode value of the respective participants scale

27 or subscale. Missing values at 18 months in the single leg stance and fast walking speed were replaced

1	with respectively four and two values from the previous test at 6 months. Due to missing data that were
2	not imputable, the number of participants in some analyses differed from sample size. Level of
3	significance was set at p-value ≤ 0.05 . Statistical analyses were performed with SPSS version 17.0 for
4	Windows.
5	

6 Descriptive statistics were used to examine demographic and test data. A large ceiling effect was found in 7 trials of the Single Leg Stance with eyes open, where about 70 % of the participants reached the upper 8 time limit. We found no floor or ceiling effect in the Single Leg Stance - eyes closed, which is thus 9 presented. Distributions of scores were examined by Q-Q plots, by comparing mean, trimmed mean and 10 median of the scales and subscales, and tests of normality. The scales are of ordinal nature, so all 11 assumptions of normality were not met. Differences within and between groups were estimated by 12 Wilcoxon Signed Ranks Test, Mann-Whitney test, and Kruskal-Wallis test in non-parametric data (43), 13 and paired and independent t-tests, and ANOVA in parametric data (44).

14

15Test-retest reliability was explored in this study group at initial evaluation. Two subsequent trials in the16single leg stance and walking speeds were used. Relative reliability was calculated by intra-class17correlation coefficients (ICC (1.1 and ICC (1.3)). Findings of the single leg stance –eyes closed showed r18> 0.7, and walking speeds showed r > 0.9. Both were considered satisfactory according to proposals (45),19p. 234. An indication of absolute reliability was calculated by within-subject standard deviation (Sw) (46,2047), Smallest detectable changes (SDC) were calculated.

21

22 Change is defined as scores at the initial test minus scores at 18 months. Standardized effect size of

23 change (ES) was used to help interpret the magnitude of change. The effect size was calculated by paired

24 t-tests, $ES = \sqrt{t^2/t^2}$ + degrees of freedom in parametric data (44), and by Wilcoxon Signed Ranks tests in

25 non-parametric data, $ES = Z / \sqrt{n}$, where n denotes the total number of non-zero observations in the

- analyses) (43). Change scores are assumed to be normally distributed when n > 20 (48). Pearson's
- 27 product moment correlation was used to examine associations between change scores. Guidelines

28 proposed by Cohen (49) were used to evaluate the strength of correlations, and effect sizes. Low

29 correlation (or small effect) is indicated by values 0.10 - 0.29; moderate correlation (or medium effect) by

30 values 0.30 - 0.49; and high correlation (or large effect) by values 0.50 - 1.0.

1

2 **Results**

At initial evaluation the majority of participants had scores in outcome measures that indicated moderate to severe impairment in perceived dizziness, and moderate to severe limitations in activities and restrictions in participation according to the outcome measures, as shown in Table 3 and Table 4. Scores in the Vertigo Symptom Scale indicated that the participants had more trouble in the subscale "vertigo/balance" than in the subscale "autonomic/anxiety" (data not shown).

9

10 Changes in outcome measures at 18 months.

11 Statistical significant change in self-perceived symptoms (impairment) by the Vertigo 12 Symptom Scale – short form at 18 months showed large effect size of difference (Table 13 3). The changes assumed to indicate improvement in body function were mainly in the 14 subscale "vertigo/balance" (data not presented). The mean score of the Vertigo 15 Symptom Scale – short form had turned below the cut-off points of "severe dizziness" 16 (Table 3). Thirty-seven persons (54%) still reported "dizziness" at 18 months, of whom 17 21 (28%) reported severe dizziness (Table 4). Changes in the Dynamic Visual Acuity 18 test, also assumed to indicate body function, improved by a median of two lines, just 19 indicating a real change and large effects size of difference (Table 3). Thirty-one 20 participants (46%) still had persisting problems with seeing clearly while moving their 21 head (Table 4).

22

Findings in the Patient Specific Functional Scale, and the Disability Scale, assumed to
indicate activity and participation, showed improvements with large effect sizes of
difference (Table 3). The improvements in the Patient Specific Functional Scale were

1	found in physical activity and sports, work related activities, social and cultural life,
2	transportation, and domestic work. Seven persons (10%) still reported severe limitations
3	in personal important activities (Table 4). The median score of the Disability Scale had
4	improved to "mild disability" (Table 3). We found that 11(16%) participants still had
5	"severe disability" (Table 4). In tests assumed to indicate changes in activity, we found
6	statistical significant improvement in the Single Leg Stance - eyes closed, and median
7	effect size of difference (Table 3). Fifty-three persons (78%) did not reach the expected
8	standing time of 30 seconds (data not shown). The change scores in walking speeds
9	exceeded measurement error, but did not reach statistical significance. Low and medium
10	effects sizes of difference were found (Table 3).
11	
12	Findings of change in the composite measure, the Dizziness Handicap Inventory,
13	showed statistical significant change. We found mean improvements with large effect
14	size of difference (Table 3) assumed to indicate body function, activity and participation
15	The mean score indicated "mild" impact of dizziness on quality of life. Five participants
16	(7%) still had "severe" impact of dizziness on quality of life (Table 4).
17	
18	In subgroup analyses, females showed significantly larger improvements in most self-
19	report measures compared to men. No significant differences were found in subgroup
20	analyses according to age, months since onset of dizziness, intervention, or between
21	vestibular and non- vestibular dizziness.
22	
23	Associations between change scores of different outcome measures.
24	We found high correlation between the Vertigo Symptom Scale assumed to indicate
25	hade for stign / impairments and the composite measure Dissinger Handison Leventery

25 body function/impairments and the composite measure Dizziness Handicap Inventory

1	(Table 5). Significant moderate and high correlation were also found between change
2	scores in the Vertigo Symptom Scale and measures assumed to indicate activity and
3	participation, the Patient Specific Functional Scale, and the Disability Scale (Table 5).
4	We found no correlation between change scores assumed to indicate body
5	function/impairments: the Vertigo Symptom Scale and the Dynamic Visual Acuity test
6	(Table 5). We found moderate correlation between the Dynamic Visual Acuity and two
7	tests assumed to indicate activity/limitations, the Single leg Stance-eyes closed and
8	preferred walking speed. No correlation was found between changes in the Vertigo
9	Symptom Scale and tests that were assumed to indicate activity: the Single Leg Stance,
10	and walking speeds (Table 5).
11	
12	Correlations between changes in outcome measures assumed to indicate activity and
13	participation showed moderate and high correlations ($r = 0.3 - 0.6$) (Table 5). The
14	highest correlations were found between the Disability Scale and Dizziness Handicap
15	Inventory, and between the Disability scale and the Patient Specific Functional Scale
16	(Table 5). A high correlation was found between preferred and fast walking speed, also
17	indicating activity. Generally, we found no correlations between self-reported measures,
18	and tests (Table 5).
19	
20	Discussion
21	In this follow-up study of persons with dizziness we found correlations between
22	change-scores at 18 months ranging $r = 0$ - 0.6. Significant correlations were found
23	between change scores of body function and activity/participation ($0.3 \le r \ge 5$). We
24	found no completions between self report measures and tests. We found the first is if and

- 24 found no correlations between self-report measures and tests. We found significant
- 25 improvements in impairments indicated by outcome measures of dizziness and visual

1	acuity, and in activity and participation indicated by outcome measures of standing				
2	balance, patient specific activities, disability, and quality of life. Similar patterns of				
3	change were also found in subgroup analyses except in gender.				
4 5 6	<i>Methodological issues</i> Number of participants lost to follow-up could represent a problem. However, no				
7	differences at initial testing, and the given reasons for withdrawals, led to the				
8	assumption that the withdrawals had minor influences on the results. The sample, only				
9	including adult up to 65 years old (mean age 46.7 years), represent results from adults,				
10	not elderly, in line with some other follow-up studies (11, 12, 14, 50-52). Since				
11	dizziness in this sample had lasted \geq 4 months, the results might be generalizable to				
12	adult patients with long lasting dizziness, but not elderly patients.				
13					
14	Symptoms and signs of dizziness may indicate disorders/disruptions of many origins.				
15	This study included persons with different types of dizziness. Investigations used to				
16	determine the origin of dizziness might have varied, but diagnostics was not the focus of				
17	this study. Vestibular origins (65%) were most frequent in this study group. Subgroup				
18	analyses showed no statistical significant differences between diagnostic groups.				
19	Physiotherapy to patients with dizziness is based on symptoms, signs, and function, not				
20	necessarily diagnoses. Persons with the same diagnosis still present differences in				
21	functioning. Previous and more recent long-term follow-ups have included participants				
22	with different origin of dizziness (8, 11, 20, 21) as we did, as well as determined				
23	specific diagnostic groups (14, 18, 19, 51). The mixed origin of dizziness in our study				
24	could be perceived as a limitation, but could also be seen as being representative of				
25	patients seen in primary health care.				
26					

1 With a small sample size there might be a risk of both Type I and Type II errors. The 2 null hypothesis of no difference between scores at initial evaluation and 18 months 3 follow-up required a significance level of ≤ 0.05 . Analyses performed in the total group 4 (n=68) are assumed to be sufficient to test the 0-hypothesis. The additional subgroup 5 analyses resulting in smaller sample sizes could be more vulnerable to errors. Analyses 6 of change-scores in samples ≥ 20 is assumed to be sufficient according to experts (48).

7

8 To explore data from outcome measures we used different statistical methods. The 9 results are evaluated in line with proposals by Marquis et al (53): by established 10 measurement properties, magnitude of change by standardized effect size of difference, 11 comparisons with known clinical change, previously determined cut-off points, and 12 known-group references. By this we intended to illuminate limitations and strengths of 13 the applied outcome measures. The varying information on measurement properties in 14 several of the applied scales and tests, is a known challenge in evaluating change in 15 general (16), and hence also in persons with dizziness (54). This might affect the 16 internal validity of the results. Awareness of the strengths and limitations in existing 17 outcome measures is necessary to interpret change. There is a need to promote further 18 explorations of measurement properties in outcome measures in this patient group.

19

By informing about follow-up assessments, testing, and test results, as well as the
benefits of physical activity at each evaluation session, the study might be seen as an
intervention in itself. These factors probably influenced results in directions of
improvements for all participants. Knowledge of assumed benefits and the fact that you
are to be assessed again increases awareness. Additionally, information about your

condition might reduce fear of symptoms, and stimulate to increase the level of general
 activity.

3

4 The same trained physiotherapist (first author) interviewed, tested, collected and plotted 5 data, which might constitute a possible bias. Being one assessor might, however, also 6 have enabled higher reliability in assessments. The data registrations were performed 7 twice allowing for comparisons and corrections, and data analyses were checked, and 8 rechecked. The intention of the study was to explore changes in persons with dizziness 9 in different domains of functioning by use of different outcome measures. The research 10 interest did not favor any specific results. We consider the possible influence on results 11 to be minimal. 12 13 Main results 14 In this long-term follow-up study by self-reports and physical tests in several domains 15 of functioning concurrently, we found associations between change scores that were 16 comparable to previous findings of associations between measures of body function and 17 structure, and activity and participation (16). None of the questionnaires or tests in the 18 present study seemed redundant, since the highest association was r = 0.6. Hence, the 19 associations indicated that the outcome measures capture different domains of 20 functioning and dimensions of change (55). 21 22 Improvements in body function indicated by self-perceived severity of symptoms (the 23 Vertigo Symptom Scale – short form) showed no association to change scores in the test 24 assumed to indicate body function, the Dynamic Visual Acuity test, or tests assumed to 25 indicate activity as the Single Leg Stance and walking speeds. Symptoms of dizziness

are assumed to reflect impairment somewhere in the vestibular system, and the physical

1 tests are assumed to provoke symptoms or reflect problems experienced by persons with 2 symptoms of dizziness. Based on this, one could expect that changes in perceived 3 symptoms of dizziness and the physical tests indicating impairment and activity would 4 correlate. The lack of correlations found support that the different outcome measures 5 address and capture different dimensions of change. Further, the findings support the 6 need to apply both self-reports of symptoms and physical tests concurrently. Our 7 findings are in line with previous findings, where results from vestibular function or 8 balance tests do not differ between groups reporting improvement versus no 9 improvement in dizziness (18, 19, 50). 10 11 Moderate correlations between improvements in body function indicated by dynamic visual acuity, and activity indicated by standing balance and preferred walking speed, could indicate central compensation in some persons, which enabled improvements in activity. About half of the participants still had remaining problems in the tests. This

12 13 14 15 could be due to sequelae after vestibular disorders (54). The non-computerized Dynamic 16 Visual Acuity test appeared to describe a relevant problem experienced by patients, as 17 well as having the ability to capture change. Although significant, the change did not 18 exceed the expected number of lines previously found following vestibular 19 rehabilitation (40). The single leg stance appeared suitable to provide information about 20 change in balance. It addresses a relevant functional problem experienced by persons 21 with dizziness, and deserves attention in exercises and patient education to prevent falls. 22 The non-significant changes in preferred walking speed could be explained by a 23 conscious choice of a safer speed. The results were found to be within the 95% 24 confidence interval of normal range according to a later review (42), and all 25 participants were able to walk at speeds ranging from 0.71 to 1.38 m/s required to cross

street intersections in large cities (56). Knowledge about clinical important change in
 walking speeds could be useful to guide interpretations of change. Comprehensive
 measures of functional balance with elements of dual task should be included in follow ups of this patient group.

5

6 The moderate and high correlations found between change scores of body function (Vertigo Symptom Scale) and activity/participation (Patient Specific Functional Scale, 7 8 Disability Scale and the Dizziness Handicap Inventory) are comparable to previous 9 findings of associations between these domains of function (16). The changes found 10 could indicate improvement in the person's ability to cope with perceived symptoms. 11 Managing symptoms and coping strategies are considered an important part of treatment 12 of patients with chronic conditions (57). Improvements found might be a result of 13 'response shift', a result of modifications of expectations and adjustments of standards 14 and change in values (58), as well as bodily adjustments and use of alternative 15 strategies. Awareness is needed in evaluating the scores, since they might also reflect 16 persisting problems with activities that have actually been dropped by the patient.

17

The use of the Patient Specific Functional Scale is to our knowledge reported for the first time in persons with dizziness. The scale appears to provide important information to the clinician, is highly relevant to the patients, and might support motivation in coping with dizziness. A person-oriented measure might contribute to determine patient oriented goals to benefit management. Exploration of measurement properties in this patient group is warranted.

24

1	The proportion of persons with persisting dizziness (impairment) at follow-up was high					
2	(53%), but within the range found in previous studies. The proportion with severe					
3	dizziness (28%), is in line with findings in some studies in persons with vestibular					
4	disorders (12, 19, 24, 59), but lower than findings at a balance clinic (11). We found					
5	few indications of "autonomic/anxiety", and psychiatric co-morbidity in the patient					
6	group, contrary to previous long term follow-ups in persons with persisting dizziness					
7	(10, 12, 59). The 21 persons reporting severe dizziness at follow-up included more					
8	persons than the proportion who reported severe limitations in activities and					
9	participation (≤ 11 persons), and fewer persons than the proportion who still had					
10	abnormal test results (\geq 32 persons). These findings at an individual level might further					
11	promote inclusion of broad assessments in these patients to capture the complexities of					
12	change in chronic dizziness.					

13

14 This study was performed before the ICF core set for patients with vertigo, dizziness 15 and balance disorders was designed (60). Our assumptions about outcome measures in 16 patients with dizziness find support in the proposed core set. We welcome the work 17 performed by this group (60), and await the following work related to developing 18 outcome measures that are relevant for this patient group in different domains of 19 functioning.

20

21 Conclusion

22 Measuring change in patients with long lasting/chronic conditions is a complex issue. 23 Comprehensive use of outcome measures in line with the ICF, indicating changes in 24 body function, activity and participation, appeared to provide information of changes in 25 different domains of functioning. To enable broad and meaningful follow-up of patients

- 1 with long lasting dizziness, generic and condition specific measures, self-reports and
- 2 tests in these domains of functioning could be further explored.

3

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8

9 **Conflicts of interest:** none to declare.

10

11 **References**

Neuhauser HK, Radtke A, von Brevern M, Lezius F, Feldmann M, Lempert T.
 Burden of Dizziness and Vertigo in the Community. Arch Intern Med.

14 2008;168(19):2118-25.

15 2. Hannaford PC, Simpson JA, Bisset AF, Davis A, McKerrow W, Mills R. The
prevalence of ear, nose and throat problems in the community: results from a national
cross-sectional postal survey in Scotland. Fam Pract. 2005;22(3):227-33.

18 3. Tamber AL, Bruusgaard D. Self reported faintness or dizziness - comorbidity
19 and use of medicines. An epidemiological study. Scand J Public Health. 2009;37:61320.

Wiltink J, Tschan R, Michal M, Subic-Wrana C, Eckhardt-Henn A, Dieterich M,
 et al. Dizziness: Anxiety, health care utilization and health behavior - Results from a
 representative German community survey. J Psychosom Res. 2009;66:417-24.

24 5. van der Windt DAWM, Dunn KM, Spies-Dorgelo MN, Mallen CD,

Blankenstein AH, Stalman WAB. Impact of physical symptoms on perceived health in
the community. J Psychosom Res. 2008;64:265-74.

- Gassmann K, Rupprecht R. Dizziness in an older community dwelling
 population: a multifactorial syndrome. J Nutr Health Aging. 2009;March 13(3):278-82.
- 29 7. Bronstein AM, Golding JF, Gresty MA, Mandala M, Nuti D, Shetye A, et al.
- 30 The social impact of dizziness in London and Siena. J Neurol. 2010;257(2):183-90.
- 8. Nazareth I, Yardley L, Owen N, Luxon LM. Outcome of symptoms of dizziness
 in a general practise community sample. Fam Pract. 1999;16(6):616-8.
- 33 9. Skoien AK, Wilhelmsen K, Gjesdal S. Occupational disability caused by
 34 dizziness and vertigo: a register-based prospective study. Br J Gen Pract.
 35 2008;58(554):619-23.
- 36 10. Kroenke K, Lucas CA, Rosenberg ML, Scherokman BJ, Herbers JE. One-year
 37 outcome for patients with chief complaint of dizziness. J Gen Intern Med. 1994;9
 38 (12):684-9.
- 39 11. Wilhelmsen K, Ljunggren AE, Goplen FK, Eide GE, Nordahl SHG. Long-term

40 symptoms in dizzy patients examined in a university clinic. BMC Ear, Nose and Throat
41 Disord. 2009;9(2):1-9.

1 12. Godemann F, Siefert K, Hantschke Bg, Neu P, Seidl R, Str"hle A. What 2 accounts for vertigo one year after neuritis vestibularis - anxiety or a dysfunctional 3 vestibular organ? J Psych Res. 2005;39:529-34. 4 13. Brandt T, Huppert T, Zingler VC, Dieterich M, Strupp M. Long-term course and 5 relapses of vestibular and balance disorders. Restor Neurol Neurosci. 2010;28(1):69-82. Best C, Eckhardt-Henn A, Tschan R, Dieterich M. Psychiatric morbidity and 6 14. 7 comorbidity in different vestibular vertigo syndromes. Results of a prospective 8 longitudinal study over one year. J Neurol. 2009(256):58-65. 9 WHO. International Classification of Functioning, Disability and Health. 15. 10 Geneva: World Health Organization; 2001. 11 Finch E, Brooks D, Stratford PW, Mayo NE. Physical Rehabilitation Outcome 16. 12 Measures. A Guide to Enhanced Clinical Decision Making. 2 ed. Hamilton, Ontario: 13 B.C. Decker Inc; 2002. 14 Whitney SL, Herdman SJ, Cohen HS, Clendaniel RA, Shumway-Cook A, Tusa 17. 15 RJ, et al. Rehabilitation Assessment and Management. In: Herdman SJ, editor. 16 Vestibular Rehabilitation. Philadelphia, PA: F.A. Davies Company; 2000. p. 331-571. 17 18. Kammerlind AS, Ledin TEA, Skargren EIB, Odkvist LM. Long-term follow-up 18 after acute unilateral vestibular loss and comparison between subjects with and without 19 remaining symptoms. Acta Oto-Laryngologica. 2005;125 946-53. 20 Zingler VC, Weintz E, Jahn K, Mike AH, D., Rettinger N, Brandt T, et al. 19. 21 Follow-up of vestibular function in bilateral vestibulopathy. J Neurol Neurosurg 22 Psychiatry. 2007;79:284-8. 23 20. Smith-Wheelock M, Shepard NT, Telian SA. Long-term effects for treatment of 24 balance dysfunction: Utilizing a home exercise approach. Seminars in hearing. 25 1991;12(3):297-302. 26 21. Cass SP, Borello-France D, Furman JM. Functional outcome of vestibular 27 rehabilitation in patients with abnormal sensory-organization testing. Am J Otol. 1996;17(4):581-94. 28 29 22. Hansson EE, Mansson NO, Hakonsson A. Falls among dizzy patients in primary 30 healthcare: an intervention study with control group. Int J Rehabil Res. 2008;31(1):51-31 7. 32 23. Cowand JL, Wrisley DM, Walker M, Strasnick B, Jacobson JT. Efficacy of vestibular rehabilitation. Otolaryngol Head Neck Surg. 1998;118(1):49-54. 33 34 Gillespie MB, Minor LB. Prognosis in bilateral vestibular hypofunction. 24. 35 Laryngoscope. 1999;109(1):35-41. Krebs DE, Gill-Body KM, Parker SW, Ramirez JV, Wernick-Robinson M. 36 25. 37 Vestibular rehabilitation: useful but not universally so. Otolaryngol Head Neck Surg. 38 2003;128(2):240-50. 39 26. Vereeck L, Wuyts FL, Truijen S, De Valck C, Van de Heyning PH. The effect of 40 early customized vestibular rehabilitation on balance after acoustic neuroma resection. 41 Clin Rehabil. 2008:22(8):698-713. 42 27. Seale J, Barnard S. Measuring and measurement tools in research. Therapy 43 research: processes and practicalities. Oxford: Butterworth-Heinemann; 1998. p. 83-92. 44 Yardley L, Burgneay J, Andersson G, Owen N, Nazareth I, Luxon LM. 28. 45 Feasability and effectiveness of providing vestibular rehabilitation for dizzy patients in the community. Clin Otolaryngol. 1998(23):422-48. 46 47 Wilhelmsen K, Strand LI, Nordahl SHG, Eide GE, Ljunggren AE. Psychometric 29. 48 properties of the Vertigo Symptom Scale - Short Form. BMC Ear, Nose and Throat 49 Disord [Internet]. 2008 cited 2008 August 17; 8(2). Available from:

50 http://www.biomedcentral.com/1472-6815/8/2

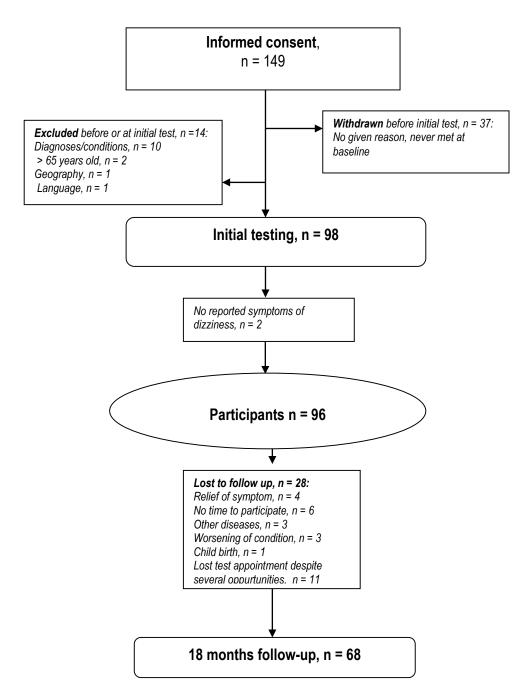
1 30. Yardley L, Donovan-Hall M, Smith H, Wash BM, Mullee M, Bronstein AM. 2 Effectiveness of primary care-based vestibular rehabilitation for chronic dizziness. Ann 3 Intern Med. 2004;141:598-605. 4 31. helsedirektoratet S-o. ICF: International klassifikasjon av funksjon, 5 funksjonshemming og helse. Trondheim Aktietrykkeriet; 2003. Stratford P, Gill C, Westaway M, Binkley J. Assessing disability and change on 6 32. 7 individual patients: A report of a patient specific measure. Physiother Can. 8 1995;47(4):258-63. 9 Shepard NT, Telian SA. Practical management of the balance disorder patient. 33. 10 London: Singular Publishing Group Inc; 1996. 11 Hall CD, Herdman SJ. Reliability of clinical measures used to assess patients 34. 12 with peripheral vestibular disorders. J Neurol Phys Ther. 2006;30(2):74-81. 13 35. Jacobson GP, Newman CW. Assessing Dizziness-Related Quality of Life. In: 14 Jacobson GP, Shepard NT, editors, Balance Function assessment and management, San 15 Diego, CA: Plural Publishing, Inc; 2008. p. 99-131. 16 Tamber AL, Wilhelmsen K, Strand LI. Measurement properties of the Dizziness 36. 17 Handicap Inventory by cross-sectional and longitudinal design. Health Qual Life 18 Outcomes [Internet]. 2009 2010 January 12; 7(101). Available from: 19 http://www.hglo.com/content/7/1/101. 20 Terwee CB, Dekker FW, Wiersinga WM, Prummel MF, Bossuyt PMM. On 37. 21 assessing responsiveness of health-related quality of life instruments: Guidelines for 22 instrument evaluation. Qual Life Res. 2003;12:349-62. 23 38. Whitney SL, Furman JM. Vestibular Rehabilitation. In: Jacobson GP, Shepard 24 NT, editors. Balance function assessment and management. San Diego, CA: Plural 25 Publishing; 2008. p. 543-83. 26 39. Whitney SL, Herdman SJ. Physical Therapy Assessment of Vestibular 27 Hypofunction. In: Herdman SJ, editor. Vestibular Rehabilitation 3ed. Philadelphia: F.A. 28 Davis Company; 2007. p. 272-99. 29 40. Badke MB, Shea TA, Miedaner JA, Grove CR. Outcomes after rehabilitation for 30 adults with balance dysfunction. Arch Phys Med Rehabil. 2004;85:227-33. 31 Bohannon RW. Comfortable and maximum walking speed of adults aged 20-79 41. 32 years: reference values and determinants. Age Ageing. 1997;26(1):15-9. 33 42. Bohannon RW, Andrews AW. Normal walking speed: a descriptive meta-34 analysis. Physiotherapy. 2011;97:182-9. 35 Field A. Non-parametric tests. Discovering statistics using SPSS. 2 ed. London: 43. 36 SAGE; 2005. p. 521-70. 37 Field A. Comparing two means. Discovering statistics using SPSS. London: 44. 38 SAGE; 2005. p. 269-308. Domholdt E. Physical Therapy Research. Principles and applications. 2 ed. 39 45. 40 Philadelphia: W.B. Saunders Company; 2000. 41 Beaton DE. Understanding the relevance of measured change through studies of 46. 42 responsiveness. SPINE. 2000;25 (24):3192-9. 43 Terwee CB, Bot SDM, de Boer MR, van der Windt DAWM, Knol DL, Dekker 47. 44 J, et al. Quality criteria were proposed for measurement properties of health status 45 questionnaires. J Clin Epidemiol. 2007;60:34-42. Altman DG. Comparing groups - continuous data. Practical statistics for 46 48. 47 medical research. London: Chapman & Hall/CRC; 1991. p. 179-228. 48 49. Cohen J. Statistical power analysis for behavioural sciences 2ed. Hillsdale, N. J.:

49 Lawrence Erlbaum 1988.

- 50. Huppert D, Strupp M, Rettinger N, Hecht J, Brandt T. Phobic postural vertigo. A
 long -term follow-up (5-15 years) of 106 patients. J Neurol. 2005;252 564-9.
- Tschan R, Best C, Beutel ME, Knebel A, Wiltink J, Dieterich M, et al. Patients'
 psychological well-being and resilient coping protect from secondary somatoform
- vertigo and dizziness (SVD) 1 year after vestibulary disease. J Neurol. 2011;258:10412.
- 52. Kammerlind AS. Recovery after acute unilateral vestibular loss and predictors
 for remaining symptoms. Am J Otolaryngol. 2011;32(5):366-75.
- 9 53. Marquis P, Chassany O, Abetz L. A comprehensive strategy for the
- 10 interpretation of quality-of-life data based on existing methods. Value Health.

11 2004;7(1):93-104.

- 12 54. Herdman SJ. Vestibular Rehabilitation 3ed. Philadelphia: F. A. Davis Company;
 13 2007 2007.
- Streiner DL, Norman GR. Health Measurement Scales a practical guide to their
 development and use. 3 ed. Oxford: Oxford University Press; 2003.
- 16 56. Robinett CS, Vondran MA. Functional ambulation velocity and distance
- 17 requirements in rural and urban communities. A clinical report. Phys Ther.
- 18 1988;68:1371-3.
- 19 57. Taylor SE. Management of Chronic Illness. Health Psychology. 7 ed. Boston:
 20 McGraw-Hill; 2009. p. 270-98.
- 21 58. Yardley L, Dibb B. Assessing subjective change in chronic illness: An
- examination of response shift in health related and goal-oriented subjective status.
 Psychol Health. 2007;22(7):813-28.
- Synthesis 24
 Synthesis 25
 Synthesis 25
 Synthesis 26
 Synthesis 27
 Synthesis 27
 Synthesis 27
 Synthesis 28
 Synthesis 28<
- 26 60. Grill E, Bronstein AM, Furman JM, Zee DS, Müller M. International
- 27 Classification of Functioning, Disability and Health (ICF) Core Set for patients with
- 28 vertigo, dizziness and balance disorders. Journal of Vestibular Research: Equilibrium &
- 29 Orientation. 2012;22(5-6):261-71.
- 30



Figur 1. Flowchart of participants and withdrawals.

Tables 1

2 3 4

Table 1 – Description of particin

Characteristics	Total group (n = 68)
Gender: female: n (% within group)	47 (69)
Age in years: mean (SD), min-max	46.7 (12.1), 26-64
Number of months since onset of symptoms: median (min-max)	21.5 (4 - 406)
Work status at present: Employed persons: n (%), Employed on sick-leave: n (%) Long term social security benefit: n (%) Others n (%)	58 (85) 21 (31) 7 (10) 3 (4)
Number of months of sick-leave/ social security benefit due to dizziness: Median (min-max)	6.0 (0 - 45)
Number of conditions/diseases: mean (SD), min-max	1.6 (1.2), 0-5
Number of medicines used: mean (SD), min-max	1.3 (1.7), 0-9

Table 2 - Number of cases of vestibular, non-vestibular and unknown origin of dizziness (n=68)

1

	Origins of dizziness		No of cases (%)
Vestibular	*Benign Paroxysmal Positional	Vertigo	6
	Dysfunction balance system		3
Dysfunction vestibular nerve NOS		1	
	Labyrinthitis		4
Mb Meniere		1	
Sacculus disorder		1	
	Vestibular dysfunction NOS		1
	Vestibular neuronitis: unilateral		19
	Vestibular neuronitis: bilateral Vestibular undecided		3 5
		Total no.	5 44 (65)
		rotarno.	++ (00)
Non vestibular	Dizziness (excludet vestibular)		1
	Dizziness NOS		2
	Dizziness syndrome NOS		1
	Dizziness and imbalance NOS		1
	Epileptic attacks		1
	Mal DeBarquement		1
	Musculoskeletal disorders		5
	Sudden deafness		1
	Traumatic Brain Injury		1
	Vertigo NOS		1
		Total no.	15 (22)
Unknown Total	NOS	Total no.	9 (13) 68 (100)

⁶ 7 8 9 10

*One patient had Benign Paroxysmal Positional Vertigo and vestibular neuronitis, only counted once NOS = No Other Specification

11

Table 3– Scores in outcome measures at initial evaluation and change scores at follow-up (N = 68) 1

					2
		Initial scores	Difference at 18 months: change scores °	Effect Si differenc	
VSS –sf t	total ^a				4
PSFS a	Mean (SD)	14.5 (9.6),	4.6 (7.1)	0.5 ^d	5
Media DS ª	an (min, max)	6.0 (0, 10)	1.7 (- 4, 9.67)	0.7 e	6
	an (min, max)	3.0 (0, 5)	1.0 (- 3, 4)	0.7 ^e	7
DVAT a	Mean (SD)	38.4 (18.6),	11.6 (15.6)	0.6 ^d	8
	an (min, max) R ^b	5.0 (0, 9)	2.0 (- 6, 6)	0.5 ^e	9
	Mean (SD)	10.4 (9.1),	- 3.0 (10.4)	0.3 ^d	10
PWS ^b	Mean (SD)	1.3 (0.3),	- 0.1 (0.2)	0.2 ^d	11
FWS [♭]	Mean (SD)	2.1 (0.3),	0.1 (0.3)	0.3 ^d	12
	. ,		· ·		13

^a Positive scores indicate improvement ;

^b Negative scores indicate improvement;

^c **Bold scores** indicate significant change, p-value < 0.05;

^d Effect size of difference calculated from t-value in paired sample t-tests;

e Effect size of difference calculated from Z value in Wilcoxon Signed Ranks test

VSS-sf = Vertigo Symptom Scale- short form;

PSFS = Patient Specific Functional Scale;

DS = Disability Scale;

DHI = Dizziness Handicap Inventory;

14 15 16 17 18 20 21 22 23 24 25 26 27 DVAT = Dynamic Visual Acuity Test; SLS-EC-R = Single Leg Stance, Eyes closed, Right leg;

PWS = Preferred Walking Speed m/s;

FWS = Fast Walking Speed m/s

- Table 4 Distribution of scores according to severity in outcome 1
- 2 measures at inclusion and follow-up (N=68).
- 3

		Initial test	Follow-up	
Measures	Categories	n (%)*	n (%)	
VSS-sf:				
'Dizziness'	\leq 11 points	30 (45)	46 (68)	
'Severe dizziness'	≥ 12 points	37 (54)	21 (28)	
	n (%)	67 (99)	67 (96)	
PSFS:				
As before	0 points	11 (16)	16 (24)	
Small limitations	1 - 3 points	8 (12)	29 (43)	
Moderate limitations	4 - 6 points	22 (32)	15 (22)	
Severe limitations	7 - 10 points	27 (40)	7 (10)	
	n (%)	68 (100)	67 (99)	
DS:				
No disability	0-1 points	18 (27)	38 (56)	
Mild and moderate	2-3 points	30 (44)	19 (28) 11 (16)	
Severe disability	4-5 points	20 (29)		
	n (%)	68 (100)	68 (100)	
DHI:				
Mild	0-30 points	26 (38)	41 (60)	
Moderate	31-60 points	31 (46)	19 (28)	
Severe	61-100 points	10 (15)	5 (7)	
	n (%)	67 (99)	64 (96)	
DVAT	≤ 3 lines	14 (21)	32 (47)	
	≥ 4 lines	50 (74)	31 (46)	
	n (%)	64 (94)	63 (93)	

VSS-sf = Vertigo Symptoms Scale – short form; PSFS = Patient Specific Functional Scale; DS = Disability Scale;

DHI = Dizziness Handicap Inventory; DVAT = Dynamic Visual Acuity test.

1 2 Table 5 – Associations between change scores in different outcome measures

(Pearson's correlation coefficient: r) (N = 68).

3

Outcome Measures:	Vertigo Symptom Scale – short form	Patient Specific Functional Scale	Disability Scale	Dizziness Handicap Inventory	Dynamic Visual Acuity test	Single leg stance - right - eyes closed	Preferred walking speed	Fast walking speed	
Vertigo Symptom Scale – short form	1								
Patient Specific Functional Scale	0.4 **	1							
Disability Scale	0.5 ***	0.5 ***	1						
Dizziness Handicap Inventory	0.5 ***	0.3 *	0.6 ***	1					
Dynamic Visual Acuity test	- 0.1	- 0.02	- 0.02	0.2	1				
Single leg stance –right – eyes closed	0.004	- 0.2	0.01	- 0.02	- 0.3 *	1			
Preferred walking speed	- 0.1	- 0.1	- 0.1	- 0.1	- 0.3 *	0.2	1		
Fast walking speed	- 0.1	0.02	- 0.1	- 0.03	0.1	- 0.1	0.5 ***	1	

4 5 **Bold scores** indicate significant change * = p-value ≤ 0.05 ; ** = p-value < 0.01; *** = p-value < 0.001

6