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Study protocol

Effects and feasibility of a standardised orientation and mobility training in using an identification cane for older adults with low vision: design of a randomised controlled trial

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Abstract

Background: Orientation and mobility training (O&M-training) in using an identification cane, also called symbol cane, is provided to people with low vision to facilitate independent participation in the community. In The Netherlands this training is mainly practice-based because a standardised and validly evaluated O&M-training in using the identification cane is lacking. Recently a standardised O&M-training in using the identification cane was developed. This training consists of two face-to-face sessions and one telephone session during which, in addition to usual care, the client's needs regarding mobility are prioritised, and cognitive restructuring techniques, action planning and contracting are applied to facilitate the use of the cane. This paper presents the design of a randomised controlled trial aimed to evaluate this standardised O&M-training in using the identification cane in older adults with low vision.

Methods/design: A parallel group randomised controlled trial was designed to compare the standardised O&M-training with usual care, i.e. the O&M-training commonly provided by the mobility trainer. Community-dwelling older people who ask for support at a rehabilitation centre for people with visual impairment and who are likely to receive an O&M-training in using the identification cane are included in the trial (N = 190). The primary outcomes of the effect evaluation are ADL self care and visual functioning with respect to distance activities and mobility. Secondary outcomes include quality of life, feelings of anxiety, symptoms of depression, fear of falling, and falls history. Data for the effect evaluation are collected by means of telephone interviews at baseline, and at 5 and 17 weeks after the start of the O&M-training. In addition to an effect evaluation, a process evaluation to study the feasibility of the O&M-training is carried out.

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This is an Open Access article distributed under the terms of the Creative Commons Attribution License (http://creativecommons.org/licenses/by/2.0), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.
**Discussion:** The screening procedure for eligible participants started in November 2007 and will continue until October 2009. Preliminary findings regarding the evaluation are expected in the course of 2010. If the standardised O&M-training is more effective than the current O&M-training or, in case of equal effectiveness, is considered more feasible, the training will be embedded in the Dutch national instruction for mobility trainers.

**Trial registration:** ClinicalTrials.gov NCT00946062

**Background**
Low vision is common among older people and is associated with mobility problems, limitations in physical and social functioning, ADL and IADL disability, reduced well-being, falls, depression, loneliness and mortality e.g. [1-15]. Particularly the level of mobility is essential for conducting activities in daily life, but this domain of functioning is directly affected by visual impairments [11,12,15,16]. Mobility problems – as a result of visual impairments – may therefore threaten independent functioning of older persons.

Mobility problems in older persons are related to a series of negative outcomes. Limitations in mobility are found to be associated with lower levels of quality of life, depression, cognitive dysfunctioning, fair or poor health, hip fracture, disability, institutionalisation and mortality [7,17-22]. Mobility limitations as a result of visual impairments may particularly limit a person’s ability to travel independently and, consequently, many aspects of daily routine life as well as life satisfaction are affected substantially [11,23-27]. For participating in social and physical activities, abilities for travelling are crucial. Travel in the environment involves skills of orientation and mobility. Orientation is the ability to recognize the environment and establish position in relation to the environment, whereas mobility is the physical ability to move in an orderly, efficient, and safe manner through the environment [28]. To maintain travel independence, it is essential for a visually impaired person to learn new orientation and mobility skills to compensate for reduced visual information.

Orientation and mobility training (O&M-training), which is a component of the rehabilitation facilities for visually impaired people, aims to maintain independence of travel by teaching visually impaired persons to ambulate and negotiate the environment safely and independently. It may decrease mobility limitations and contribute positively to societal participation and quality of life. O&M-training is often supplemented by the use of assistive devices [29]. Common mobility devices that have been used in O&M-training among persons with visual impairments are canes, such as the identification cane and the long cane [28]. With training in the use of visual and non-visual information including assistive devices, visually impaired persons gain a better understanding of their environment, which enables them to travel more comfortably, efficiently, and safely. Early intervention might be essential to slow down the progression of disablement [30], as persons with visual impairment are particularly at risk of developing disability [29,31,32]. Although the benefits of O&M-training have been object of study, the conducted evaluation studies have substantial limitations such as the lack of a control group [33,34] or randomisation [28], or the inclusion of rather small study populations [28,35]. Furthermore, none of these studies focussed on other societal outcomes next to mobility such as (social) participation and quality of life. Assessing multiple outcomes in effect studies are, however, recommended since Crews and colleagues indicated that interventions in the area of visual impairments may have effects on one kind of outcomes, such as mobility, but not on others, such as quality of life [36].

In The Netherlands, no previous research on the impact of O&M-training on daily functioning in older people with visual impairments has been conducted. In addition, evaluation studies of O&M-training in the use of an identification cane, also called symbol cane, which is frequently part of rehabilitation care for adults with low vision in The Netherlands, were hardly identified in the international literature. And finally, although there is a Dutch national instruction on orientation and mobility for mobility trainers employed in low vision rehabilitation care, there is no standardised protocol in delivering such O&M-training to older adults with low vision.

**Objectives of the study and the current paper**
The main objectives of the study are:

1) to develop a standardised O&M-training in the use of an identification cane by older adults with low vision;

2) to evaluate this newly developed standardised training with respect to effectiveness and feasibility in a randomised controlled trial.

The current paper presents the design of the randomised controlled trial that will evaluate the standardised O&M-training in the use of the identification cane by older adults with low vision in The Netherlands.
Methods/Design

Study design and setting

A parallel group randomised controlled trial was developed to evaluate a standardised O&M-training in the use of an identification cane (NCT00946062). The trial is conducted in collaboration with the two main organisations for low vision rehabilitation care in The Netherlands: Bartiméus and the VisioSensisDeBrink Group (as of January 2010: Royal Visio). In 18 local centres, which are scattered over The Netherlands, O&M-training is provided to older persons with visual impairment. The centres were randomly allocated to the control group or intervention group stratified by organisation (at that time: Bartiméus – four local centres, Sensis – four local centres, and Visio – ten local centres).

To screen eligible people for the trial, the local centres provide the research team with information on older clients with low vision who may receive mobility support. After the screening procedure and baseline measurement clients in the control group receive usual care, i.e. the regular O&M-training in using the identification cane, and clients in the intervention group receive the standardised O&M-training in using the identification cane. Follow-up data for the effect evaluation are collected at 5 and 17 weeks after the start of the training by means of telephone interviews. To obtain data for the process evaluation trainers complete registration forms after each training session and clients are interviewed by telephone at 8 weeks after the start of the training. Figure 1 shows the design of the trial. The Medical Ethical Committee of the Maastricht University/Academic Hospital Maastricht granted approval for conducting this trial.

Study population

The screening of potential participants started in November 2007 and occurs in a stepped procedure. Firstly, during the local centre's standard exploratory interview a staff member of a local centre roughly screens a potential participant using a six-item pre-structured registration form. The items include whether the client: (1) suffers from low vision, (2) is 55 years of age or older, (3) lives independently or in a home for older people, (4) experiences difficulty to avoid large obstacles due to the vision loss, (5) suffers from additional impairments that cause full inability to leave the home, and (6) consents to an additional screening interview by telephone for potential participation in the trial. If items 1, 2, 3 and 6 are scored ‘yes’ and items 4 and 5 are scored ‘no’, contact information is recorded and the registration form is sent to the research team. Secondly, a pre-structured 10-minute interview by telephone is conducted by the research team to determine the client's eligibility for participation in the trial. The additional inclusion criteria are: 1) able to go outside for a short walk or doing groceries, and 2) experiencing difficulties with safely crossing a street, or experiencing difficulties with recognising acquaintances outdoors, or willing to become recognisable as partially sighted by means of the identification cane. Lastly, mobility trainers decide during an individual contact with the client whether O&M-training in the use of an identification cane is the eligible care for the client.

Clients are excluded if one of the following criteria is met: a) cognitive impairment (a score of less than 4 on the Abbreviated Mental Test 4 (AMT4) during an interview by telephone) [37,38], b) unable to complete an interview by telephone due to language or hearing problems, c) unable to participate in or finish the O&M-training due to confinement to bed or possible nursing home admission, d) permanent use of a walking aid that is incompatible with the use of an identification cane, e) having recently received an O&M-training in the use of an identification cane and permanent use of this cane outdoors, and f) not receiving an O&M-training in the use of an identification cane as treatment for their mobility problem.

Eligible clients receive oral and written information about the trial during the first and the second step of the screening procedure. Those clients who signed an informed consent form were included in the trial. An overview of all inclusion and exclusion criteria is shown in Table 1. To maintain active participation of the local centres, the staff receives regular updates with respect to the number of clients and their progress in the trial by means of newsletters, monitoring telephone contacts and weekly reviews by email.

Randomisation and masking

After completing a baseline interview by telephone and confirmation on the eligibility of the O&M-training in the use of an identification cane by the mobility trainer, clients are officially assigned to the regular or the standardised O&M-training. To prevent contamination bias randomisation stratified by organisation was performed at the level of the local centres. Randomisation at the client or mobility trainer level may lead to contamination because (1) a mobility trainer may then train clients in the control group as well as clients in the intervention group or (2) a mobility trainer may unintentionally inform other mobility trainers about the standardised O&M-training. To decide during an individual contact with the client whether O&M-training in the use of an identification cane is the eligible care for the client.

The collaborating organisations for low vision rehabilitation care in The Netherlands, i.e. Bartiméus and VisioSen-
sisDeBrink Group, consented with participation in the trial before randomisation of the local centres. To control for differences in organisation characteristics, randomisation of local centres was performed per organisation (at that time: Bartiméus, Sensis and Visio). To assure even distribution of local centres with respect to expected number of clients per month and to avoid contamination due to close collaboration of several local centres, all local centres per organisation were matched in pairs: Bartiméus – four local centres, one pair; Sensis – four local centres, two pairs, and; Visio – ten local centres, three pairs. According to a predefined format one of each pair was randomly allocated to the intervention group or control group. This was carried out by an external party who threw a dice; even numbers corresponded to allocation to the intervention group and odd numbers to allocation to the control group.

Due to the nature of the trial and the information provided to clients before the start of the trial (required by the Medical Ethical Committee), mobility trainers, researchers and participants are not masked to intervention status. Participants are, however, not informed about their intervention status until the start of the O&M-training. Trained outcome assessors who perform the interviews by telephone are masked for intervention status.
Despite the presence of a Dutch national instruction on orientation and mobility for mobility trainers employed in low vision rehabilitation care, there is no fixed protocol for O&M-training in the use of an identification cane. To obtain insight into this O&M-training as provided by the mobility trainers, information was collected between March and June 2007 by means of: 1) observing O&M-training sessions (n = 5), 2) studying the literature of the Dutch national instruction on orientation and mobility and attending its session on the identification cane, and 3) performing individual face-to-face interviews with mobility trainers (n = 18) of the organisations for rehabilitation care for people with visual impairment in The Netherlands. The interviews were guided by a questionnaire. In addition to questions related to the characteristics of the current O&M-training in the use of an identification cane (see Table 2), mobility trainers were asked to reflect on elements missing in the current O&M-training and potential opportunities to improve this training. Based on information obtained from the mobility trainers and the national instruction on orientation and mobility, a new standardised O&M-training was developed.

The standardised O&M-training aims to facilitate safe and independent participation in the community by optimal use of one's abilities, and to facilitate uptake of old or new activities. To achieve these aims several strategies were added to the regular O&M-training, that is: prioritising the client's needs, cognitive restructuring, action planning, contracting, providing direct feedback and stimulating individual problem solving, and finding personal, realistic solutions. The standardised O&M-training consists of two face-to-face sessions and one telephone session. Compared to the regular O&M-training, the standardised O&M-training is well structured, yet still tailor-made as clients are actively involved in their rehabilitation with respect to mobility. For example, they are encouraged to individualise their training by means of formulating personal goals regarding activities in the community. Table 2 presents the main characteristics of the regular and standardised O&M-training. More information with respect to the development of the standardised O&M-training will be presented elsewhere.

Mobility trainers were eligible if they provided O&M-training in the use of an identification cane to partially sighted clients of one of the organisations. Trainers who would provide the standardised O&M-training to the intervention group received a two-hour instruction by the researcher (GZ). To prepare for the instruction, trainers read the manual of the standardised O&M-training. After the instruction, mobility trainers informed the researcher if difficulties in applying the standardised O&M-training were observed or questions arose. To monitor whether participants received sufficient care, trainers answered two questions as part of the process evaluation. These questions included to what extent the O&M-training met the participant's need for mobility support according to the perception of the trainer and whether additional training sessions in the use of the identification cane were needed. Trainers were instructed to provide additional O&M-training sessions to the clients if needed. Trainers providing care according to the standardised O&M-training needed to complete both the 2 face-to-face sessions and the one telephone session before conducting extra training sessions on O&M.

Orientation and mobility training

Despite the presence of a Dutch national instruction on orientation and mobility for mobility trainers employed in low vision rehabilitation care, there is no fixed protocol for O&M-training in the use of an identification cane. To obtain insight into this O&M-training as provided by the mobility trainers, information was collected between March and June 2007 by means of: 1) observing O&M-training sessions (n = 5), 2) studying the literature of the Dutch national instruction on orientation and mobility and attending its session on the identification cane, and 3) performing individual face-to-face interviews with mobility trainers (n = 18) of the organisations for rehabilitation care for people with visual impairment in The Netherlands. The interviews were guided by a questionnaire. In addition to questions related to the characteristics of the current O&M-training in the use of an identification cane (see Table 2), mobility trainers were asked to reflect on elements missing in the current O&M-training and potential opportunities to improve this training. Based on information obtained from the mobility trainers and the national instruction on orientation and mobility, a new standardised O&M-training was developed.

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Table 1: Screening for participants: inclusion and exclusion criteria

| Inclusion criteria:                                      | Exclusion criteria:                                      |
|---------------------------------------------------------|---------------------------------------------------------|
| - Aged 55 years or over                                  | - Cognitive impairment (a score of less than 4 on the Abbreviated Mental Test 4) [37,38] |
| - Low vision                                            | - Language or hearing problems that impede completing an interview by telephone |
| - Living independently in the community or in a home for older people | - Confinement to bed or possible nursing home admission that impede completion of the O&M-training |
| - Able to see large obstacles and to go outside for a short walk or doing groceries | - Permanent use of a walking aid incompatible with the use of an identification cane |
| - One of the following:                                 | - Having recently received an O&M-training in the use of an identification cane and permanent use of this cane |
| * experiencing difficulties with safely crossing a street |                                                          |
| * experiencing difficulties with recognising acquaintances outdoors |                                                          |
| * willing to become recognisable as being partially sighted by using the identification cane |                                                          |
| - Written informed consent                               |                                                          |
| - Orientation and Mobility training (O&M-training) in the use of an identification cane |                                                          |
No restrictions were held with respect to receiving care other than O&M-training in the use of the identification cane. In The Netherlands the costs of the O&M-training are reimbursed; the costs associated with the purchase of the cane are paid by the consumers.

**Measures**

**Effect evaluation**

The concepts measured to determine the effectiveness of the standardised O&M-training comprise to a great extent the ‘Activity and Participation’ domains of the International Classification of Functioning, Disability and Health [ICF; [39]]. As ICF domains we include ‘self care’, ‘mobility’, ‘domestic life’, ‘interpersonal interactions and relationships’, and ‘community, social and civic life’, respectively. An overview of the main outcomes, the number of items, the interpretation of the scoring and the measurements during the course of the study is presented in Table 3.

The primary outcomes of the effect evaluation are ADL self care, as assessed by the 11-item ADL subscale of the Groningen Activity Restriction Scale (GARS) [40], and the subscale distance activities and mobility of the Visual Functioning Questionnaire (VFQ) [41,42]. The secondary outcomes comprise domestic, community, and social and civic life assessed by the subscales domestic, outdoor activities and leisure/work, respectively, of the Frenchay Activities Index (FAI) [43,44], and interpersonal interactions and relationships, assessed as social support interactions by the Social Support List (SSL12-I) [45] and as feelings of loneliness by a one-item question (according to Zijlstra and
colleagues [46]). Health-related quality of life is assessed by the EQ-5D that comprises 5 items on mobility, self-care, usual activities, pain/discomfort and anxiety/depression [47]. In addition, several aspects with respect to mental functioning, fear of falling and falls are assessed: mental health and dependency which is a subscale of the VFQ [41,42], feelings of anxiety and symptoms of depression as assessed by the Hospital Anxiety and Depression Scale subscales anxiety (HADS-A) [44,48] and depression (HADS-D) [44,48], concerns about falling as assessed by the 16-item Falls Efficacy Scale International (FES-I) [49,50] and a one-item scale [46], as holds as well for activity avoidance due to fear of falling [46] and falls history indoors and outdoors [46].

With respect to activities frequently present in the O&M-training, being recognised as partially sighted and the use of the identification cane, several questions were formulated (not tabulated). First, related to safety, participants are presented four activities, i.e. crossing a street safely, crossing a major intersection safely, travelling with public transportation and walking in a crowded place. For each of the four activities participants indicate their perceived skills (1 = not at all skilled to 4 = very skilled), their feelings of safety (1 = not at all safe to 4 = very safe) and the frequency in which they perform the activity (1 = never to 4 = almost daily/often). Second, participants indicate how often other people show them consideration in traffic situations and in the city centre or a shop and how often they feel ashamed of their vision loss (1 = very often/often to 4 = never). Lastly, participants indicate to what extent the identification cane is used (1 = never to 4 = almost daily/often) – and, if applicable, in which locations -, how often they expect to use the cane in the upcoming 3 months (1 = never to 4 = almost daily/often), and if applicable, the reason for not using the identification cane.

All primary and secondary outcomes of the effect evaluation were assessed at baseline, and at 5 and 17 week follow-up, except for the outcomes regarding the use of the identification cane. The latter outcomes were solely assessed at 5 and 17 week follow-up. Before randomisation data on descriptive variables and covariates were collected, i.e. with respect to a) the participants: age, gender, marital status, living situation, educational level, employment, year of becoming visually impaired, primary diagnosis, use of an assistive device for mobility purposes, presence of a chronic medical condition (5 major medical conditions were derived from a 19-item checklist [51]), and the problem for which mobility support was requested, and b) the trainers: age, gender, educational level, completion of occupational training, completion of the Dutch national orientation and mobility instruction for mobility trainers, organisation employed, and years of working experience.

### Process evaluation

To identify programme factors that may influence the effectiveness of the standardised O&M-training and factors that may contribute to future improvement and implementation of the training a process evaluation is carried out. The main outcomes of the process evaluation are

#### Table 3: Outcome measures of the effect evaluation

| Variable | No. of items | Range* |
|----------|--------------|--------|
| **Primary outcome measures** | | |
| ADL self care (ADL subscale of the GARS) [40] | 11 | 11 to 44 |
| Distance activities and mobility (subscale of the VFQ) [41,42] | 8 | 8 to 40 |
| **Secondary outcome measures** | | |
| Activities index (FAI) [43,44] | 15 | 15 to 60 |
| Social support interactions SSL12-I [45] | 12 | 12 to 48 |
| Feelings of loneliness [46] | 1 | 1 to 6 |
| Health-related quality of life (EQSD) [47] | 5 | 5 to 15 |
| Mental health and dependency (subscale of the VFQ) [41,42] | 6 | 6 to 30 |
| Feelings of anxiety (HADS-A) [44,48] | 7 | 0 to 21 |
| Symptoms of depression (HADS-D) [44,48] | 7 | 0 to 21 |
| Concerns about falling (FES-I) [49,50] | 16 | 16 to 64 |
| Fear of falling [46] | 1 | 1 to 5 |
| Activity avoidance due to fear of falling [46] | 1 | 1 to 5 |
| Number of indoor falls in the previous 6 months† [46] | 1 | 1 to 6 |
| Number of outdoor falls in the previous 6 months† [46] | 1 | 1 to 6 |

* The bold scores indicate the most favourable scores.
† At FU1 and FU2 the number of falls between the previous and the current assessment is measured.
ADL = activities of daily life; GARS = Groningen Activity Restriction Scale; VFQ = Visual Functioning Questionnaire; EQSD = EuroQol 5 Dimensions; HADS-A = anxiety subscale of the Hospital Anxiety and Depression Scale; HADS-D = depression subscale of the Hospital Anxiety and Depression Scale; FAI = Frenchay Activities Index; FES-I = falls efficacy scale international.
Data collection
Data for the effect evaluation are collected at baseline, and at 5 and 17 weeks after the start of the O&M-training during 40-minute interviews by telephone. Trained interviewers, who are masked for group allocation, perform the interviews. In addition, a 25-minute interview by telephone is performed at 8 weeks after the start of the O&M-training to obtain data for the process evaluation from participants in the intervention group as well as the control group. Pre-structured questionnaires on process aspects are used to gather data from the mobility trainers per participant. Trainers receive this questionnaire before the first session of the participant’s O&M-training. As recommended by Hollis and Campbell [55], non-compliant participants are approached for all follow-up assessments.

Sample size and power
The power calculation is based on data from Kempen and colleagues [40] with regard to the primary outcome variable: GARS ADL self care. Two times 67 participants will provide 80% power at alpha .05 to detect differences between the intervention and control groups’ mean change score of at least 3.5 points (SD is 8.1 equivalent with an effect size of .43 on the GARS ADL self care). However, a dropout during the study of 25% may be expected. Therefore, 2 × 95 participants are needed for randomisation.

Analysis
Data of the effect evaluation will be analysed according to intention-to-treat principle and on-treatment principle. Univariate techniques and mixed-effects regression analyses will be applied to test for between-group differences with respect to the primary and secondary outcome measures at all follow-up assessments. Models will be adjusted for relevant covariates, including age, gender and baseline differences. Descriptive techniques will be used to describe participants of the trial and to analyse the data of the process evaluation.

Discussion
O&M-training is a common health care service provided by organisations for rehabilitation care for people with visual impairment. The training aims to maintain independence of travel by teaching visually impaired persons to ambulate and negotiate the environment safely and independently. It may decrease mobility limitations and contribute positively to societal participation and quality of life. In an ageing population it is important to positively influence these domains to prevent physical, mental and social dysfunctioning which can lead to disability and institutionalisation. Yet, evidence on the effects of O&M-training in older adults with visual impairment is scarce, particularly with respect to the O&M-training in the use of an identification cane. This paper presents the design of a randomised controlled trial that will evaluate the effects and feasibility of a standardised O&M-training in the use of the identification cane by older adults with low vision in The Netherlands.

Progress of the study
The screening procedure for eligible participants started in November 2007 and will continue until October 2009. In April 2009 28 participants (14 control group, 14 intervention group) had completed the training and follow-up assessments and another 11 participants (7 control group and 4 intervention group) had finished the O&M-training and the follow-up assessment at 5 weeks.

Recruitment of the participants by the collaborating organisations Bartiméus and VisioSensisDeBrink Group is slower than expected based on a dossier study by Verstraten and colleagues [56] and information obtained by the first author during interviews with mobility trainers. Reasons for the slow recruitment of participants as stated by the organisations include, among others, lack of eligible clients, lack of interest of clients to participate in the study, other research studies target the same population, work load of the mobility trainers and employees who perform the screening, unwillingness of mobility trainers to wait for the necessary research actions – such as obtaining informed consent and performing an baseline interview – before starting the training, adoption of new computerization system, etc. Although numerous actions have been undertaken to facilitate the organisations in the process of screening and to limit the time for research actions, recruitment of sufficient participants remains the main challenge of this study. The financial budget for the study allows to screen for eligible participants until October 2009. The 2 × 95 participants based on the power calculation will not be enrolled at that time.

Detailed information regarding the development of the standardised O&M-training and the effect and process evaluation will be presented in forthcoming papers. Preliminary findings regarding the evaluation are expected in the course of 2010.
Table 4: Outcome measures of the process evaluation

| Variable                                                                 | Measurement |
|--------------------------------------------------------------------------|-------------|
| **Population reached**                                                   |             |
| Target population and proportion of the intended target population       | TI-Rp, SQt  |
| General characteristics of the participants and trainers                | Qt          |
| Number of participants that refused, dropped out or completed the training| Qt          |
| Reasons for withdrawal                                                  | Qt          |
| **Performance according to protocol**                                   |             |
| Format, preparation time and duration of the session                     | Qt          |
| Per session component: extent to which carried out, duration and active participation by the participant* | Qt          |
| Extent to which the trainer achieved to:                                 |             |
| - related to identification cane: provide information, raise the participant’s awareness of its advantages, demonstrate the use and have the participant experience the use of the cane* | Qt          |
| - have the participant phrase his/her important activities related to mobility and how to perform activities safely and independently* | Qt          |
| - have the participant setting goals regarding an action plan*           | Qt          |
| - teach orientation skills* and teach mobility skills*                  | Qt          |
| **Exposure and engagement**                                             |             |
| Total number of sessions                                                | Qt          |
| Use of materials*                                                       | TIp, Qt     |
| Overall opinion of the trainer/participant regarding the participant’s engagement in: |             |
| - the training                                                          | TIp, Qt     |
| - formulating of an action plan and carrying out an action plan*         | Qt          |
| Exposure and adherence to homework                                      | TIp         |
| Extent to which the participant complied with contracts*                | Qt          |
| Quality of actions plans formulated by the participants*                | Qt          |
| **Opinion on the training**                                             |             |
| Overall opinion on the training                                         | TIp, Qt     |
| Opinion regarding:                                                     |             |
| - number, duration and course of the sessions                           | TIp, Qt     |
| - homework and comprehensibility of the complete training               | TIp         |
| - number of extra training sessions needed                              | TIp, Qt     |
| - whether the O&M-training met the participant’s need for mobility support | TIp, Qt     |
| Usefulness regarding:                                                  |             |
| - discussing mobility problems, receiving information on different kinds of canes and practicing with the identification cane | TIp         |
| - formulating an action plan and carrying out an action plan*           | TIp         |
| - contracting with respect to formulating and carrying out action plans* | TIp         |
| Burden experienced by the participant                                   | TIp         |
| Recommendation of the training to others                                | TIp         |
| Overall opinion on the trainer                                          | TIp, Qt     |
| **Perceived benefit or achievement**                                    |             |
| Benefit regarding:                                                     |             |
| - the O&M-training                                                      | TIp         |
| - advantages of an identification cane and self-confidence              | Qt          |
| - realistic action planning*                                            | Qt          |
| Use of identification cane in daily life                                | Qt          |
| Achievement regarding:                                                 |             |
| - identification cane: knowledge, aware of advantages and having experienced the cane | TIp         |
| - able to think how to perform an activity safely and independently      | TIp         |
| - self-confidence and performance of an activity safely and independently| TIp         |
| - formulating a realistic action plan* and carrying out an action plan* | TIp         |
| - skills for orientation and skills for mobility                        | TIp         |
**Table 4: Outcome measures of the process evaluation (Continued)**

| Deviations of each session component* | Qt | Tlp, Qt |
|---------------------------------------|----|--------|
| Main goal of contact - time spent on issues not related to O&M-training | Qt | Tlp, Qt |
| Strong and weak aspects of the O&M-training | Qt | Tlp, Qt |
| Hampering and stimulating factors for use of the standardised O&M-training* | Qt | Tlp, Qt |
| Matters for improvement - materials and standardised O&M-training* | Qt | Tlp, Qt |

*Assessed in the intervention group only.
O&M-training = orientation and mobility training; TI-Rp = telephone interview prior to randomisation with the participant (as part of the screening process); SQt = starting questionnaire, i.e. questionnaire filled in by the trainer before training his/her first participant of the study; Qt = questionnaire filled in by the mobility trainer per participant; Tlp = interview by telephone with the participant.

**Future implementation**
Throughout the development of the standardised O&M-training, future implementation in the Dutch health care for visually impaired people has been taken into account. An implementation and dissemination plan is designed. One of the main actions consists of embedding the standardised O&M-training in the Dutch national instruction for mobility trainers, i.e. if the standardised O&M-training is more effective than the current O&M-training or, in case of equal effectiveness, is considered more feasible.

**Abbreviations**
O&M: training refers to orientation and mobility training; ADL: activities of daily living; IADL: instrumental activities of daily living; ICF: refers to the International Classification of Functioning, Disability and Health.

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

**Authors’ contributions**
GZ, GK, GvR, ES, DB, JvdV and PV contributed to the concept and design of the study. GZ drafted the manuscript with input from the other authors. All authors read and approved the final manuscript.

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**References**
1. Berger S, Porell F: The association between low vision and function. J Aging Health 2008, 20(5):504-525.
2. Brody BL, Gamst AC, Williams RA, Smith AR, Lau PW, Dolinak D, Rapaport MH, Kaplan RM, Brown SL: Depression, visual acuity, comorbidity, and disability associated with age-related macular degeneration. Ophthalmology 2001, 108(10):1893-1900.
3. Broman AT, Munoz B, Rodriguez J, Sanchez R, Quigley HA, Klein R, Snyder R, West SK: The impact of visual impairment and eye disease on vision-related quality of life in a Mexican-Ameri- can population: proyecto VER. Invest Ophthalmol Vis Sci 2002, 43(11):3393-3398.
4. de Boer MR, Plijim SM, Lips P, Moll AC, Volker-Dieben HJ, Deeg DJ, van Rens GH: Different aspects of visual impairment as risk factors for falls and fractures in older men and women. J Bone Miner Res 2004, 19(9):1529-1547.
5. Evans JR, Fletcher AE, Wormald RP, Ng ES, Stirling S, Smeeth L, Breeze E, Bullpit CJ, Nunes M, Jones D, et al: Prevalence of visual impairment in people aged 75 years and older in Britain: results from the MRC trial of assessment and management of older people in the community. Br J Ophthalmol 2002, 86(7):795-800.
6. Jacobs JM, Hammerman-Rozenberg R, Maaravi Y, Cohen A, Stessman J: The impact of visual impairment on health, function and mortality. Aging Clin Exp Res 2005, 17(4):281-286.
7. Kempen GI, Verbrugge LM, Merrill SS, Ormel J: The impact of multiple impairments on disability in community-dwelling older people. Age Ageing 1998, 27(5):595-604.
8. Koski K, Luukinen H, Laippala P, Kivela SL: Risk factors for major injurious falls among the home-dwelling elderly by functional abilities. A prospective population-based study. Gerontology 1998, 44(4):232-238.
9. Limburg H: Epidemiologie van visuele beperkingen en een demografische verkenning. Een studie in opdracht van Sticht- ting InZicht. Grootebroek: Health Information Services; 2007.
10. Mojan-Azzi SM, Sousa-Poza A, Mojan DS: Impact of low vision on well-being in 10 European countries. Ophthalmologica 2008, 222(3):205-212.
11. Montarzino A, Robertson B, Aspinall P, Ambrecht A, Findlay C, Hine J, Dhillon B: The impact of mobility and public transport on the independence of visually impaired people. Visual Impair Res 2007, 9:67-82.
12. Owsley C, McGwin G Jr: Association between visual attention and mobility in older adults. J Am Geriatr Soc 2004, 52(11):1901-1906.
13. Paz SH, Globe DR, Wu J, Azen SP, Varma R: Relationship between self-reported depression and self-reported visual function in Latinos. Arch Ophthalmol 2003, 121(7):1021-1027.
14. Sakari-Rantala R, Era P, Rantanen T, Heikkinen E: Associations of sensory-motor functions with poor mobility in 75- and 80-year-old people. Scand J Rehabil Med 1998, 30(2):121-127.
15. Salive ME, Guralnik J, Glynn RJ, Christen W, Wallace RB, Ostfeld AM: Association of visual impairment with mobility and physical function. J Am Geriatr Soc 1994, 42(3):287-292.
16. Maino JH: Visual deficits and mobility. Evaluation and management. Clin Geriatr Med 1996, 12(4):803-823.

17. Borghin G, Jakobsson U, Edberg AK, Hallberg IR: Self-reported health complaints and their prediction of overall and health-related quality of life among elderly people. J Nurs Stud 2005, 42(2):147-158.

18. Guralnik JM, Simonsick EM, Ferrucci L, Glynn RJ, Berkman LF, Blazer DG, Scherr PA, Wallace RB: A short physical performance battery assessing lower extremity function: association with self-reported disability and prediction of mortality and nursing home admission. J Gerontol 1994, 49(2):H185-94.

19. Lampinen F, Heikkinen E: Reduced mobility and physical activity and a prediction of depressive symptoms among community-dwelling older adults: an eight-year follow-up study. Aging Clin Exp Res 2003, 15(3):205-211.

20. Melzer D, Lan TY, Guralnik JM: The predictive validity for mortality of the index of mobility-related limitation – results from the ERF study. Age Ageing 2003, 32(6):619-625.

21. Shumway-Cook A, Ciol MA, Yorkston KM, Hoffman JM, Chan L: Use of assistive devices in daily activities and visual impairment for visually impaired persons over age 60. J Orthop Sports Phys Ther 1998, 28(3):143-160.

22. Soong GP, Lovie-Kitchin JE, Brown B: Visual deficits and mobility. Evaluation and management. Clin Geriatr Med 1996, 12(4):803-823.

23. Wahl HW, Heyl V, Schilling O: The validity of the Hospital Anxiety and Depression Scale. An updated literature review. J Psychosom Res 2002, 52(2):69-77.

24. yardley L, Beyer N, Hauer K, Kempen GI, Piot-Ziegler C, Todd C: Development and initial validation of the Falls Efficacy Scale International (FES-I). Age Ageing 2005, 34(6):614-619.

25. Zijlstra GA, van Haastregt JC, van Eijk JT, Kempen GI: Pre-publication history for a group of working-age adults. Optom Vis Sci 2007, 84(8):775-784.

26. Holbrook M, Skillbeck CE: An activities index for use with stroke patients. Age Ageing 1983, 12(2):166-170.

27. Langelaan M, van Nispen RM, Knol DL, Moll AC, de Boer MR, Wouters B, van Rens G: Visual Functioning Questionnaire: revalidation of psychometric properties for a group of working-age adults. Optom Vis Sci 2008, 85(9):1601-1610.

28. Wahl HW, Heijl V, Stahl JW: A program in orientation and mobility: experiences of visually impaired older people. Br J Occup Ther 2008, 71(10):414-421.

29. Wahl HW, Heijl V, Schilling O: The role of vision impairment for the outdoor activity and life satisfaction of older adults: a process evaluation. Visual Impair Blin 1999, 7(4):143-160.

30. Soong GP, Lovie-Kitchin J, Brown B: Does mobility performance of visually impaired adults improve immediately after orientation and mobility training? Optom Vis Sci 2001, 78(9):567-566.

31. Dahlin-Ivanoff S, Sonn U: Use of assistive devices in daily activities among 85-year-olds living at home focusing especially on the visually impaired. Disabil Rehabil 2004, 26(24):1423-1430.

32. Geruschat DR, Turano KA, Stahl JW: The validity of the Hospital Anxiety and Depression Scale. An updated literature review. J Psychosom Res 2002, 52(2):69-77.

33. Verbrugge LM, Jette AM: The disablement process. Soc Sci Med 1994, 38(1):1-14.

34. Cacciarelli F, Abete P, Maggi S, Luchetti G, Calabrese C, Vlto L, Leoza D, Ferrara N, Vitale DF, Rengo F: Disability and 6-year mortality in elderly population. Role of visual impairment. Aging Clin Exp Res 2004, 16(5):382-388.

35. Ivanoff SD, Sonn U, Lundgren-Lindqvist B, Sjostrand J, Steen B: Disability in daily life activities and visual impairment: a population study of 85-year-old people living at home. Scand J Occup Ther 2000, 7(4):148-155.

36. Geruschat DR, Turano KA, Stahl JW: Traditional measures of mobility performance and retnits pigmenteda. Optom Vis Sci 1998, 75(7):523-537.

37. Straw LB, Hart RK, Zimmerman GJ: A program in orientation and mobility for visually impaired persons over age 60. J Visual Impair Blin 1991, 85(3):108-113.

38. Johnson JT, Johnson BF, Blash BB, de l’Aune WD: Gait and long cane kinematics: a comparison of sighted and visually impaired subjects. J Orthop Sports Phys Ther 1998, 27(2):162-166.

39. Crews JE, Long RG: Conceptual and methodological issues in rehabilitation outcomes for adults who are visually impaired. J Visual Impair Blin 1997, 91(2):135-147.

40. Crews JE, Nightingale PG: Evaluation of a shortened version of the Abbreviated Mental Test in a series of elderly patients. Clin Rehab 1997, 11(3):243-248.

41. Swain DG, O’Brien AG, Nightingale PG: Cognitive assessment in elderly patients admitted to hospital: the relationship between the shortened version of the Abbreviated Mental Test and the Abbreviated Mental Test and Mini-Mental State Examination. Clin Rehabil 2000, 14(6):608-610.

42. WHO: International Classification of Functioning, Disability and Health (ICF). Geneva: World Health Organization; 2001.

43. Kempen GIJM, Miedema I, Ormel J, Molenaar W: The assessment of disability with the Groningen Activity Restriction Scale: Conceptual framework and psychometric properties. Soc Sci Med 1996, 43(11):1601-1610.

44. Stelmack JA, Stelmack TR, Massof RW: Measuring low-vision rehabilitation outcomes with the NEI VFQ.25. Invest Ophthalmol Vis Sci 2002, 43(9):2859-2868.

45. Langelaan M, van Nispen RM, Knol DL, Moll AC, de Boer MR, Wouters B, van Rens G: Visual Functioning Questionnaire: revalidation of psychometric properties for a group of working-age adults. Optom Vis Sci 2007, 84(8):775-784.

46. Holbrook M, Skillbeck CE: An activities index for use with stroke patients. Age Ageing 1983, 12(2):166-170.

47. CBS, Jong, de Haan R, Li, M, Groenier KH: The Frenchay Activities Index. Assessment of functional status in stroke patients. Stroke 1993, 24(8):1173-1177.

48. Kempen GIJM, Van Eijk LM: The psychometric properties of the SSLI 12-I, a short scale for measuring social support in the elderly. Soc Indic Res 1995, 35(3):303-312.

49. Rabin R, de Chorro F: EQ-5D: a measure of health status from the EuroQol Group. Ann Med 2001, 33(5):337-343.

50. Bjelland I, Dahl AA, Haug TT, Neckelmann D: The validity of the Hospital Anxiety and Depression Scale. An updated literature review. J Psychosom Res 2002, 52(2):69-77.

51. Yardley L, Beyer N, Hauer K, Kempen GI, Piot-Ziegler C, Todd C: Process evaluation of a shortened version of the Falls Efficacy Scale International (FES-I). Age Ageing 2005, 34(6):614-619.

52. Kempen GIJM, Todd CJ, van Haastregt JC, Zijlstra GAR, Beyer N, Freiburger E, Hauer K, Piot-Ziegler C, Yardley L: Cross-cultural validation of the Falls Efficacy Scale International (FES-I) in older people: results from Germany, the Netherlands and the UK were satisfactory. Disabil Rehabil 2007, 29(2):155-162.

53. Jonkers C, Lamers F, Bosma H, Metsemakers J, Kempen G, Van Eijk J: Process evaluation of a minimal psychological intervention to reduce depression in chronically ill elderly persons. Patient Educ Couns 2007, 68(3):252-257.

54. van Haastregt JC, Zijlstra GA, van Rossum E, van Eijk JT, Kempen GI: Pre-publication history for a group of working-age adults. Optom Vis Sci 2008, 85(8):775-784.

55. Verstraten P, Oudshoorn J, van Grinsven R: Mobilitieitsvragen van blinde en slechtziende ouderen. Dossierstudie naar hulpvra- gen en interventies. Grave. Sensis; 2006.

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