Social participation in working-age adults with aphasia: An updated systematic review

by

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Abstract

**Background:** A previous systematic review found limited data regarding social participation in working-age people with aphasia (PWA). This population has many roles to fulfill, that are negatively affected by aphasia. A review of recent studies may reveal more information on the challenges in re-establishing social roles and thus may inform treatment thereof.

**Method:** The aim was to provide an updated systematic review on social participation in PWA under 65 years of age. Studies from 2005-2017 were searched from Scopus, Pubmed and Psychinfo. Search terms were derived from the International Classification of Functioning, Disability and Health (ICF) and the Aphasia-Framework for Outcomes Measures (A-FROM). Aspects of domestic life, interpersonal relations and interactions, education and employment and community, civic and social life were investigated.

**Results:** From 2,864 initial hits, 11 studies were identified, all of which were on the American Speech-Language-Hearing Association (ASHA) Level III of evidence. The studies indicated that participation in domestic life is reduced and PWA showed reduced social networks, loss of friendships and changes in the quality of marital relations. Few PWA returned to work or spent time on education. Limitations in community, civic and social life were noted and there were contradictory findings on the impact of contextual factors on social participation. There was an increase in research into contextual factors impacting on social participation in PWA and in the use of conceptual frameworks in the last decade.

**Conclusions:** Social participation in working-age adults is limited across the social domains. While the ICF conceptual framework is increasingly used, no studies used the A-FROM. There is greater use of standardised assessments and larger sample sizes.

**Keywords:** stroke, aphasia; social participation, working-age, young, middle-age, International Classification of Functioning, Disability and Health (ICF), Aphasia-Framework for Outcomes Measures (A-FROM)
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DECLARATION

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I declare that this research report is my own original work. Where secondary material is used, this has been carefully acknowledged and referenced in accordance with university requirements.

I understand what plagiarism is and am aware of the University of Pretoria’s policy in this regard.

[Signature]

15/05/2017

SIGNATURE    DATE
Acknowledgements

Special thanks go to Professor Alta Kritzinger and Bhavani Pillay, the research supervisors, for their patience, guidance and assistance. This study would not have been possible without them.

I would like to thank my family and friends for their encouragement and support during this entire process. Lastly, unending thanks goes to my Heavenly Father.
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<tr>
<td>A-FROM</td>
<td>Aphasia- Framework for Outcome Measurement</td>
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<td>ASHA</td>
<td>American Speech-Language-Hearing Association</td>
</tr>
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<td>BMI</td>
<td>Body Mass Index</td>
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<td>CVA</td>
<td>Cerebral Vascular Accident</td>
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<td>FIM</td>
<td>Functional Independence Measure</td>
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<td>HRQL</td>
<td>Health Related Quality of Life</td>
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<tr>
<td>ICF</td>
<td>World Health Organisation International Classification of Functioning, Disability and Health</td>
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<td>LPAA</td>
<td>Life Participation Approach to Aphasia</td>
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<td>MRI</td>
<td>Magnetic Resonance Imaging</td>
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<tr>
<td>NOS</td>
<td>Newcastle-Ottawa Quality Assessment Scale</td>
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<td>NPO</td>
<td>Non-Profit Organisation</td>
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<tr>
<td>OT</td>
<td>Occupational Therapist</td>
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<tr>
<td>PRISMA-P</td>
<td>Preferred Reporting Items for Systematic Reviews and Meta-Analyses Protocols</td>
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<tr>
<td>PWA</td>
<td>Person with Aphasia</td>
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<tr>
<td>Q</td>
<td>Quantitative</td>
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<td>QL</td>
<td>Qualitative</td>
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<td>QOL</td>
<td>Quality of Life</td>
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<td>R</td>
<td>Range</td>
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<td>SD</td>
<td>Standard Deviation</td>
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CHAPTER 1

Aim of the chapter:

The aim of this chapter is to orientate the reader to the topic of aphasia in working-age adults and resulting social participation difficulties, state the research problem, present a literature overview of related studies and to conclude with the rationale, research question of the study and the terminology as used in the dissertation.

1.1. Introduction

Worldwide, stroke is included among the foremost causes of long-term disability and often results in decreased quality of life (QOL) and work ability (Bohra et al., 2015; R. J. P. Dalemans, De Witte, Wade, & Van den Heuvel, 2008). Global stroke incidence rates range from 41 to 316 per 100 000 population, per year (Thrift et al., 2014). Within the upper margin of this range is the Agincourt rural sub-district of South Africa, with an estimated crude stroke incidence of 244 per 100 000 person years\(^1\) (Maredza, Bertram & Tollman, 2015). A study conducted in the Greater Cincinnati/Northern Kentucky region indicates that there is a significant increase in stroke incidence rates between the ages of 20 and 54 years (Kissela et al., 2012).

This concerning increase in strokes in the young and middle-age population may be related to a number of factors. The prevalence of the risk factors for stroke is changing in the working-age population, with a significant increase in diabetes, high cholesterol and obesity over time (Kissela et al., 2012). Kissela et al. (2012) found a marked increase in coronary heart disease and an increased rate of drug abuses was noted over time in young individuals with stroke. Maredza, Bertram, Gómez-Olivé, and Tollman (2016) found that high blood pressure and a high Body Mass

\(^1\) Refers to the number of new cases within a population at risk, in a specified time period. The incidence proportion (number of new cases within a specified time period divided by the size of the population at risk) is divided by the number of years over which the population was studied.
Index (BMI) were associated with a significant portion of the stroke burden in rural South Africa. In addition, it may be that with the advances in technology and the use of Magnetic Resonance Imaging (MRI), a greater number of strokes are being detected in the young, who have been shown to undergo more MRIs than older patients (Kissela et al., 2012). The increase in stroke in the working-age population is of particular concern to speech-language therapists due to the frequently presenting aphasia and its wide-spread consequences.

1.2. Aphasia

One of the main manifestations of stroke is aphasia, with a frequency ranging from 21 to 38% (Croquelois & Bogousslavsky, 2011; Dickey et al., 2010; Hoffmann & Chen, 2013; Pedersen, Vinter, & Olsen, 2004). The frequency variation is due to differences in diagnostic criteria, post-stroke time period prior to evaluation as well as the selection criteria of the participants in studies (Kyrozis et al., 2009). Aphasia is an acquired neurogenic disorder that involves language difficulties that may affect reading, writing, comprehending and expressing language, not due to difficulties with sensation, motor functioning or intellect (Chapey, 2008). Aphasia is often classified into subtypes based on the presenting symptoms and the manner in which each modality is affected. These subtypes include fluent aphasias (Wernicke’s aphasia, conduction aphasia and transcortical sensory aphasia), non-fluent aphasias (Broca’s, transcortical motor and global aphasia) and anomic aphasia (Chapey, 2008). Researchers have found some discrepancy in the frequency of the presentation of the subtypes of aphasia. Global aphasia tends to be the most frequent type of aphasia identified (Kang et al., 2010; Pedersen et al., 2004). Kang et al. (2010) identify anomic aphasia as relatively on par with Broca’s aphasia in frequency, followed by anomic aphasia, while Pedersen et al. (2004) identifies Wernicke’s aphasia as occurring more frequently than Broca’s aphasia in the acute stage. Aphasias have been found to be heterogeneous in both the aetiology and subtype of aphasia (Hoffmann & Chen, 2013).

1.3. Impact of age on aphasia

It appears that age is another factor contributing to the heterogeneous nature of aphasia. Research indicates that young people with aphasia (PWA) are a unique
population. Advancing age has been shown to be associated with an increased risk of aphasia (Dickey et al., 2010; Plowman, Hentz, & Ellis, 2012). In first ischemic stroke individuals, age has been found to be a predictor for aphasia, with the risk increasing by 4% with each year (Engelter et al., 2006; Kadojić et al., 2012). The type of stroke can be classified as ischemic (blocked blood vessel) or haemorrhagic (burst blood vessel) in origin (Andersen, Olsen, Dehlendorff, & Kammersgaard, 2009). A haemorrhagic stroke occurs less frequently than an ischemic stroke (approximately ten times less frequently), but tends to be associated with a higher mortality risk or a stroke of a greater severity (Andersen et al., 2009).

Younger individuals more often present with non-fluent aphasia, while older individuals are more likely to present with fluent aphasia (Croquelois & Bogousslavsky, 2011; Eslinger & Damasio, 1981; Plowman et al., 2012). It appears that with age and possible neuropathological variation, a stroke located near the middle cerebral artery will either shift more posteriorly (leading to Wernicke’s aphasia) or cover most of the middle cerebral artery region (leading to global aphasia) (Eslinger & Damasio, 1981; Godefroy, Dubois, Debachy, Leclerc, & Kreisler, 2002). While a difference in aphasia symptoms and types is evident across ages, Godefroy et al. (2002) found that this is true for ischemic strokes only, with adults with conduction aphasia being younger and those with subcortical aphasias being older. It appears that the age-aphasia association is related to the effects of ageing on vascular pathology and therefore the lesion location (Godefroy et al., 2002).

It appears that the age difference does not have a significant impact on the symptoms of aphasia in the acute stage but rather in the chronic stage (Godefroy et al., 2002; Pedersen et al., 2004). This may be due to a number of reasons, one of which is that the ‘true’ aphasia type may be masked by initial non-fluency in the acute stage, as most patients progress from non-fluent to fluent aphasia types. Pedersen et al. (2004) also suggest that the hypothesis of the lateralisation of comprehension in the brain with increasing age and possible reductions in brain plasticity at an older age, particularly for comprehension, may result in this finding. Furthermore, older patients may have early and undiagnosed Alzheimer’s disease,
which may contribute to the receptive difficulties evident at assessment (Pedersen et al., 2004).

There is considerable controversy in the literature regarding the impact of age on prognosis in recovery in aphasia. Some research supports the theory that prognosis deteriorates with an increase in age (Ali, Lyden, & Brady, 2015; Laska, Hellblom, Murray, Kahan, & Von Arbin, 2001; Tsouli, Kyritsis, Tsagalis, Virvidaki, & Vemmos, 2009) and others do not (Liang et al., 2001; Pedersen et al., 2004). Plowman et al. (2012) conclude in their review that while age has been identified as a prognostic indicator for stroke recovery, this has not specifically been determined for aphasia or language recovery, likely due to the wide variety of factors that impact on language in an individual.

1.4. Impact of aphasia on short and long-term functional outcomes

While there are variations in the impact of age on aphasia and recovery, it is well known that aphasia, in general, is one of the strongest predictors of poor functional recovery and social outcome following a stroke (Fang et al., 2003). Aphasia may result in significant short and long-term challenges. Short-term, the presence of aphasia is associated with longer hospital stays and extended use of rehabilitation services (Dickey et al., 2010). PWA present with lower Functional Independence Measure (FIM) scores (both cognitive and motor) on admission to and discharge from hospitals or rehabilitation centres (Gialanella, Bertolinelli, Lissi, & Prometti, 2011). PWA are less likely to return home as compared to individuals with stroke, but without aphasia (Gialanella et al., 2011).

Long-term, Dalemans et al. (2008) found that working-age PWA show decreased participation in numerous life areas due to the often significant communication difficulties. Dalemans et al. (2008) found a reduction in domestic life; alteration in interactions and relationships as well as changes in roles as partner, family member, parent, friend and citizen following the onset of aphasia. PWA show a reduction in the number of social contacts and a shift from relationships with friends and family to professionals (R. J. P. Dalemans et al., 2008). PWA between the ages of 15 and 49 show decreased employment rates as opposed to adults without aphasia, but in the same age group (Naess, Hammersvik, & Skeie, 2009). Furthermore, it appears that
should the PWA return to work, it is often at a less demanding level (R. J. P. Dalemans et al., 2008). These wide-spread restrictions are of concern as research shows that activity limitations negatively impact on health related quality of life (HRQL) in PWA (Hilari, Needle, & Harrison, 2012).

1.5. The shift towards participation

In an attempt to facilitate functional outcomes in PWA, a number of social and life participation approaches have been developed. The goal for using these frameworks and approaches is to maximise re-engagement in life and to improved functional participation of PWA in the context of their individual lives and activities (Kagan & Simmons-Mackie, 2007; Roth & Worthington, 2005). One such approach is the Life Participation Approach to Aphasia (LPAA), which emphasises the concerns of the PWA within the treatment and decision-making process (Hallowell, 2017). Another framework, the World Health Organisation International Classification of Functioning, Disability and Health [ICF] (Hallowell, 2017; World Health Organisation [WHO], 2001), views health conditions according to two domains, namely ‘Functioning and disability’ and ‘Contextual factors’ (Figure 1).

Figure 1. World Health Organisation International Classification of Functioning, Disability and Health framework

This approach helps to direct the focus to health, well-being and QOL, rather than to just the primary impairment (Hallowell, 2017). The multiple independent domains,
including Body Function and Structure (one’s physiological functioning and the anatomical parts of one’s body), Activities and Participation (the execution of tasks and the involvement in a life situation) and Environment and Personal factors (personal, social and attitudinal environment), interrelate to create QOL of the individual (WHO, 2001). The condition alone does not impact the degree to which an individual participates in life; the three domains interact to facilitate or decrease participation and QOL. A conceptual guide has also been developed for aphasia that is compatible with the ICF, namely the Living with Aphasia-Framework for Outcome Measurements [A-FROM] (Kagan et al., 2008). This conceptual framework was developed to guide outcome measurement in aphasia, with emphasis on real-life outcomes. It integrates QOL and domains related to the environment, participation, and personal identity in the same framework as the impairment (Kagan et al., 2008). This conceptual guide was informed by the social model of disability, whose proponents see disability in the context of society. Poor participation is seen as a result of barriers in society and the view of both the individual and people around them that disability is a problem. The disability itself is seen as far less important than the impact of the condition on engagement in meaningful activity (Hallowell, 2017). Byng and Duchan (2005) found that the social model principles and philosophies are effective in guiding treatment in aphasia.

These approaches and models are strongly reflective of the intervention goals identified by PWA. Both PWA and their family members identify increased life participation as a key area in which they would like to improve (Isaksen, 2014). A study by Worrall et al. (2011) found that the goals identified by PWA most often fell within the ‘Activity and Participation’ domain of the ICF framework (Kagan et al., 2008). A qualitative meta-analysis investigating ‘living successfully with aphasia’ identified participation in meaningful activities as fundamental by PWA, their families and speech-language therapists (Brown, Worrall, Davidson, & Howe, 2012). Of concern is the finding that speech-language therapists continue to be anchored in traditional medical models of treatment of aphasia (Gauvreau, Le Dorze, Laliberté, & Alary Gauvreau, 2016). Medical models place emphasis on the impairment, considering the causes of the condition and the specific changes to the structure of the body. Treatment will therefore focus primarily on isolated deficits. The risk of this model is treatment addressing underlying deficits, without considering participation...
and facilitation of new skills in everyday life (Hallowell, 2017). The importance of this is seen in a study conducted in Nigeria, where it was found that community reintegration is often poor in stroke survivors (Akosile et al., 2016). The researchers emphasise that particular attention must be given to mobility, vocational and social skills, two of which items fall within ‘Activity and Participation’ on the ICF (Akosile et al., 2016).

1.6. Social participation in working-age adults

In line with the development of these approaches and the subsequent drive towards their use in assessment and treatment, there appears to be increasing research into social participation and QOL of PWA (Dalemans et al., 2008; Hilari et al., 2012). Research shows that individuals with stroke, both with and without aphasia, in young and middle adulthood face particular challenges to regain a meaningful level of involvement in society in their possible roles as providers, partners and parents, which older individuals may no longer be required to perform (R. J. P. Dalemans et al., 2008; Putaala et al., 2009). Furthermore, the stroke may contribute to a lifetime of medical complications, in addition to living an extended number of years with the effects of the stroke (Jacobs, Boden-Albala, Lin, & Sacco, 2002).

While an increased interest in participation and QOL is evident, there seem to be a number of gaps in the literature. Hilari et al. (2012) found that there is limited research into HRQL for aphasia, with most studies focusing on the stroke population overall, with few researchers isolating PWA within the population (Northcott, Moss, Harrison, & Hilari, 2016). In their comprehensive systematic review of articles published from 1970 to 2005 on social participation of PWA under the age of 65, Dalemans et al. (2008) identified a number of factors to consider. While it is clear that all the domains of social participation are affected in young PWA, the researchers advise caution in the interpretation of their findings (Dalemans et al., 2008). The review identified the need for an increase in the use of a clear conceptual framework and well-defined concepts in order to better interpret the literature (Dalemans et al., 2008). They found that many of the studies investigated included small sample sizes and a limited number of standardised assessments. The researchers were unable to identify studies describing community, civic and social life, one of the domains in ‘Activity and Participation’ of the ICF framework, as an
important part of the studies. They note a lack of research into the impact of the environment on social participation and on return to work, which are key considerations in the working-age population. It is evident that much is still unknown about the participation of young and middle-aged adults with aphasia.

1.7. Rationale

The extensive review by Dalemans et al. (2008) included research studies until 2005, more than a decade past. When considering this time lapse, the clear recommendations of the authors for further research, as well as an increasing incidence of stroke in the young population, an update of recent publications and the quality thereof is required to inform current practice in speech-language therapy. With this knowledge, both assessment and treatment procedures can be adjusted to the unique needs of the working-age population for improved functional outcomes. These considerations lead to the research question: What is known about the impact of aphasia on social participation in working-age adults with stroke-related aphasia and what is the level of evidence of these studies from the last decade?

1.8. Terminology as used in the dissertation

Social participation

Social participation is defined as the ‘actual performance of activities in social life domains in interactions with others in the context in which they live (R. J. P. Dalemans et al., 2008)’. ‘Activity and Participation’ is one of the domains identified on the ICF framework, along with Body Structure and Functions and Environmental and Personal factors (WHO, 2001). Participation refers to involvement in various life situations, which on the ICF includes a number of domains: domestic life, interpersonal relations and interactions, education and employment (major life areas) and community, civic and social life (WHO, 2001). An individual may experience certain participation restrictions as a result of a condition that is impacting on their body structure and function. The individual’s participation in these domains is affected by contextual factors, including their environment and personal factors (WHO, 2001).

Working-age adults
Working-age adults have been defined as individuals from the age of 18 years (the average age of completion of secondary school) to 65 years of age, which is the common age of retirement. This period of time is one in which most individuals are engaged in some form of gainful employment. The purpose of this study is to investigate the impact of aphasia on this population that is still performing a wide variety of life roles, including bringing in an income.
CHAPTER 2

Aim of the chapter:
The aim of this chapter is to state the purpose of the research study as well as to describe the design, study criteria and selection as well as the data extraction and risk of bias assessment procedures. This chapter elaborates on the methods used in the systematic review as the restrictions on the article length do not allow for comprehensive descriptions.

2.1. Aim

The aim of this research project is to provide an updated systematic review on the social participation of working-age adults with aphasia in the last decade (2005-2017), with emphasis on the level of evidence of the research included.

2.2. Study design

A systematic review was completed to investigate the proposed research question. Systematic reviews form the basis for developing practice guidelines and they provide information on gaps in the literature and therefore inform future research (Shamseer et al., 2015). Close attention was given to the level of evidence and risk of bias in each of the studies, as the strength of evidence on a specific topic determines whether intervention guidelines can be formulated with confidence. Dalemans et al. (2008) conducted a systematic review on social participation in the working-age population with aphasia, and included studies until 2005. The authors highlighted a number of gaps in the literature and the need for further research, particularly with more consistent use of a conceptual framework (R. J. P. Dalemans et al., 2008). With the development of the ICF framework in 2001, it was presumed that a greater number of studies in this field would make use of this framework, allowing for greater consistency in the concepts used as well as address domains of participation regarding which little is known. These factors highlighted the need for an updated systematic review including studies from 2005 to the present. The Preferred Reporting Items for Systematic Reviews and Meta-Analyses Protocols
[PRISMA-P] (Shamseer et al., 2015) checklist was used to guide and structure the review (Appendix A). The PRISMA-P is a 27 numbered item checklist to guide protocol development of systematic reviews and meta-analyses evaluating therapeutic efficacy, however it is also recommended for any form of systematic review, qualitative or quantitative, because of the general lack of existing protocol guidance (Shamseer et al., 2015). The checklist contributed towards ensuring the comprehensiveness and transparency of this review on working-age adults with aphasia. As the study did not involve human participants directly, no institutional ethical clearance was required to conduct the research.

2.3. Study inclusion criteria

Articles pertaining to the various domains of social participation in working-age adults with aphasia were included in the study. The following electronic databases were searched, based on the relevance to the subject field and in line with item 10 on the PRISMA-P checklist: Scopus, PsychINFO and PubMed (Shamseer et al., 2015). The main search terms were: ‘aphasia’, and/or ‘stroke’, together with the following terms related to social participation according to the ICF (WHO, 2001) and A-FROM (Hallowell, 2017) frameworks: domestic life, relationships, education, employment, leisure, community life, social life, civic life. Additional search terms included ‘quality of life’, ‘long-term outcomes’, ‘well-being’ and ‘self-esteem’. These same search terms used in the Dalemans et al. (2008) study were used to ensure comprehensiveness of the search as well as to provide comparable findings.

2.4. Study selection

A total of three searches were conducted across the three databases with the last search being run on 24/01/2017. Using this electronic search strategy (limited to 2005-2017, English only and only original articles and reviews), the initial search yielded 2,864 articles. The specified time limit was set as the review completed by Dalemans et al. (2008) only included studies until 2005. The eligibility criteria were developed according to item eight on the PRISMA-P checklist (Shamseer et al., 2015). An electronic software program, Covidence (https://www.covidence.org), was used in order to synthesise searches from the three databases as well as to identify duplicates and to review abstracts and full-text articles. Following the removal of
duplicates, 1,625 articles remained. Systematic reviews were excluded to make use of only original studies. The identified articles' titles and abstracts were reviewed. Articles that discussed aphasia in conjunction with any of the identified social participation aspects or search terms were included for full-text review. Forty-one articles meeting the criteria were identified for full-text review.

A number of exclusion criteria were identified for the final review phase:

- Fewer than six PWA between the ages of 18 and 65
  - This criterion was identified to eliminate single case studies and to ensure that the working-age population was sufficiently represented in the study. Case studies represent the lowest level of evidence on the American Speech-Language Hearing Association (ASHA) evidence rating scale (ASHA, 2004), and may not significantly contribute to a systematic review.

- The mean age of the aphasia participants was more than 65
  - In an attempt for the research findings to reflect the performance of working-age adults, the mean age of the participants had to be below 65 years

- Quantitative studies in which the percentage of participants with aphasia was less than 10%.
  - This aspect was identified to ensure the aphasia population was adequately represented in the study.

- Participants with aphasia were not separately outlined in the study population characteristics.
  - In order to determine whether the participants met the inclusion criteria, the PWA had to be separately described in order for studies to be comparable.

- PWA that were not stroke-related
  - Studies in which aphasia was as a result of various conditions e.g. Traumatic Brain Injury were excluded to provide findings that were applicable to stroke-related aphasia and to ensure homogeneity of the study population.

- Participants presenting with aphasia after more than one stroke
An individual with more than one stroke may have different impairments from each incident. These impairments may influence findings in the assessment. The results may not purely be related to aphasia, but may be affected by a concomitant condition from a previous stroke.

- Only the acute phase (0-3 months) was reported on
  - In the acute phase, PWA are frequently recovering medically and participating in intensive rehabilitation programs. In order to obtain a true reflection of the individual's social participation, he/she had to no longer be in the acute phase but rather have already returned home and be engaged in his/her daily life.

- The measurement instrument, methodology and/or statistical/qualitative analysis were not described
  - It is not possible to compare studies or interpret the findings accurately if the instruments, methodology and data analysis are not well described.

- Spouses or carers described their own needs in relation to the PWA.
  - While this is an interesting focus of research, the current study was investigating social participation of the PWA.

- QOL was measured by an assessment tool, but findings were not described in terms of social participation
  - A number of studies investigated both QOL and HRQL of PWA. Many, however, simply measured these aspects using an assessment tool, but the findings were not described in terms of the domains of social participation, often simply obtaining a final score for each participant.

- Social participation aspects were not separately described in the study outcomes
  - It is not possible to analyse study results if there is no specific discussion regarding one or more of the aspects that fall within the social participation domains.

- Changes in QOL or social participation following a treatment program or approach.
This aspect is again an interesting topic to investigate, however the aim of the current study was to describe social participation in relation to aphasia, and not the impact of various protocols on the participation of this group.

Two of the researchers independently reviewed the 41 articles. These articles included both quantitative and qualitative studies reporting on various aspects of social participation of working-age PWA. There were conflicting decisions on seven of the articles, and following discussion between the researchers, each item was resolved. Eleven articles were included in the final review (Figure 2).

Table 1 provides a summary of the final selection of articles. Of these studies, five were qualitative in nature and six were quantitative. Six articles included aspects of domestic life, six investigated interpersonal relations and interactions and seven referred to education and employment. Only four articles included aspects of community, civic and social life and three referred to contextual factors impacting on social participation. Most of the studies were conducted in Europe, with the
exception of two, one conducted in South Korea and another in Brazil. It appears that the majority of studies were completed in middle to high income countries (World Bank Group, 2017), which may have a greater number of resources to facilitate participation, with regards to the environment and education of the population. Social participation in developing or low income countries may be negatively affected by limitations in these same factors, which might lead one to tentatively conclude that PWA may experience greater barriers to social participation in low income countries.
Table 1. Characteristics of included studies

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<tr>
<th>Author and year of publication</th>
<th>Social participation aspect/domain(s) studied</th>
<th>Q/QL</th>
<th>Country</th>
<th>Sampling method</th>
<th>Mean age in months (R/SD)</th>
<th>Sample size</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dalemans, De Witte, Wade, &amp; Van den Heuvel (2010)</td>
<td>Contextual factors impacting on social participation</td>
<td>QL</td>
<td>Netherlands</td>
<td>Purposive sampling, with criteria to include different aphasia severities, genders, levels of mobility and time post-onset</td>
<td>57.4 (R 45-71)</td>
<td>13 PWA</td>
</tr>
<tr>
<td></td>
<td>Domestic life, Education and employment, Factors impacting on social participation</td>
<td>Q</td>
<td>Netherlands</td>
<td>Purposive sampling</td>
<td>64.2 (R 35-87)</td>
<td>150 PWA</td>
</tr>
<tr>
<td>Darrigrand et al. (2011)</td>
<td>Domestic life, Factors impacting on social participation</td>
<td>Q</td>
<td>France</td>
<td>Convenience sampling, part of a larger study of stroke patients with aphasia</td>
<td>Severe aphasia: 63.7 (SD 15.1)</td>
<td>27 PWA (severe)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>Moderate aphasia: 64.1 (SD 10.4)</td>
<td></td>
</tr>
<tr>
<td>Fotiadou, Northcott, Chatzidaki, &amp; Hilari (2014)</td>
<td>Interpersonal relations and interactions, Education and employment, Community, civic and social life, Factors impacting on social participation</td>
<td>QL</td>
<td>Greece</td>
<td>Purposive sampling of blogs sustained by a sole author who had aphasia following a stroke, and which reflected on their social network</td>
<td>48.8 (R 26-69)</td>
<td>10 PWA</td>
</tr>
<tr>
<td>Hilari &amp; Northcott (2006)</td>
<td>Interpersonal relations and interactions, Education and employment</td>
<td>Q</td>
<td>United Kingdom</td>
<td>Cluster sampling framework</td>
<td>61.6 (R 21-92)</td>
<td>83 PWA</td>
</tr>
<tr>
<td>Łapkiewicz &amp; Grochmal-bach (2008)</td>
<td>Interpersonal relations and interactions</td>
<td>QL</td>
<td>Poland</td>
<td>Purposive sampling of married couples, with one partner with severe aphasia</td>
<td>58.7</td>
<td>22 PWA (and partner)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>21 Stroke, no aphasia (and partner)</td>
</tr>
<tr>
<td>Authors</td>
<td>Domains of Focus</td>
<td>Data Collection Methodology</td>
<td>Sample Characteristics</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>---------------------------------</td>
<td>-----------------------------------------------------------------------------------</td>
<td>---------------------------------------------------------------------------------------------</td>
<td>-------------------------------------------------------------</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Lee, Lee, Choi, &amp; Pyun (2015)</td>
<td>Domestic activities, Education and employment, Community, civic and social life</td>
<td>Convenience sampling of patients with post-stroke aphasia (≥6 months)</td>
<td>59.2 (SD 7.2) 32 PWA</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Matos, Jesus, &amp; Cruice (2014)</td>
<td>Domestic life, Interpersonal relations and interactions, Education and employment, Community, civic and social life</td>
<td>Unclear</td>
<td>65 (R 41-80) 14 PWA</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Mazaux, Lagadec, De Sèze, Zongo, Asselineau, Douce, Trias, Delair, Darrigrand (2013)</td>
<td>Domestic life</td>
<td>Convenience sampling, part of a larger study of stroke patients with aphasia</td>
<td>65.1 (R 13.5) 100 PWA</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Naess, Hammersvik, &amp; Skeie (2009)</td>
<td>Interpersonal relations and interactions, Education and employment</td>
<td>Purposive sampling from computer-based hospital registries</td>
<td>42 (R 24-49) 20 PWA</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pommerehn, Delboni, &amp; Fedosse (2016)</td>
<td>Domestic activities, Interpersonal relations and interactions, Community, civic and social life</td>
<td>Convenience sampling from members of a group</td>
<td>48.25 (R 25-67) 12 PWA</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Abbreviations: CVA: Cerebral Vascular Accident, NPO: Non-Profit Organisation, OT: Occupational Therapist, PWA: Person with Aphasia, Q: Quantitative, QL, Qualitative, R; Range, SD: Standard deviation, SLT: Speech-language therapist
2.5. Data management and data items

An electronic software program, Covidence, was used to synthesise searches from the three databases as well as to identify duplicates and to review abstracts and full-text articles, in accordance with item 11a of the PRISMA-P checklist (Shamseer et al., 2015). Two of the researchers independently reviewed the articles identified for full-text review. There were conflicting decisions on seven of the identified articles, and following discussion amongst the researchers, each item was resolved. Each of the selected articles was reviewed and relevant data was extracted and compiled into summaries. Once a summarised table was developed and then divided into the four social participation domains, a second review of the extracted data was completed in order to ensure accuracy. The data items investigated are as follows (Item 12 on the PRISMA-P checklist):

1. Characteristics of the study, including title, author(s), year of publication, country where the study was conducted, type of study (qualitative or quantitative), sampling method, mean age, sample size, social participation domain investigated, measurements or instruments used as well as the assessment period post-stroke

2. The impact of aphasia on the four identified social participation domains according to the ICF framework, including domestic life, interpersonal life, education and employment and community, civic and social life.

2.6. Risk of bias in selected studies

An adapted version of the Newcastle-Ottawa Quality Assessment Scale [NOS] (Wells et al., 2014) was used (Appendix B). This 'star system' tool was developed to assess the quality of non-randomised studies ultimately for incorporating the assessments in the interpretation in systematic reviews (Wells et al., 2014). Each study was judged on three broad categories: study group selection; group comparability and outcomes with a greater number of stars indicating a higher level of evidence (Wells et al., 2014). The content validity and inter-rater reliability of this rating scale have been established (Wells et al., 2014).
The widely accepted ASHA level of evidence rating scale was also used to categorise the selected studies (ASHA, 2004). This scale rates studies on four levels, with the lowest being IV and the highest being I (Table 2). Levels I and II are further differentiated into ‘a’ and ‘b’. The rating scale aids in identifying studies of a higher level of evidence to contribute to accuracy in comparison of data from numerous studies. This evidence rating was independently completed by two of the researchers, and differences noted were resolved after discussion.
<table>
<thead>
<tr>
<th>Source</th>
<th>ASHA level of evidence</th>
<th>Selection (Maximum 5 stars)</th>
<th>Comparability (Maximum 2 stars)</th>
<th>Outcome (Maximum 3 stars)</th>
<th>Total stars</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mazaux et al. (2013)</td>
<td>III</td>
<td>(b) somewhat representative of the average in the target population*</td>
<td>(a) justified and appropriate*</td>
<td>(a) controlled for aphasia*</td>
<td>8/10</td>
</tr>
<tr>
<td></td>
<td></td>
<td>1. (a) comparability between respondents’ and non-respondents’ characteristics is established and satisfactory*</td>
<td>(a) validated measurement tool on follow up**</td>
<td>1. (c) self-report</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>2. (b) hospital controls</td>
<td>(b) controlled for age, language*</td>
<td>2. N/A</td>
<td></td>
</tr>
<tr>
<td>Dalemans, de Witte, Beurskens et al. (2010)</td>
<td>III</td>
<td>(b) somewhat representative of the average in the target population*</td>
<td>(a) justified and appropriate*</td>
<td>(a) controlled for aphasia*</td>
<td>7/10</td>
</tr>
<tr>
<td></td>
<td></td>
<td>1. (a) comparability between respondents’ and non-respondents’ characteristics is established and satisfactory*</td>
<td>(a) screening tool with moderate validity*</td>
<td>1. (c) self-report</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>2. (b) hospital controls</td>
<td>(b) controlled for age, time post stroke and premorbid conditions*</td>
<td>2. Yes</td>
<td></td>
</tr>
<tr>
<td>Darrigrand et al. (2011)</td>
<td>III</td>
<td>(c) selected group of participants</td>
<td>(b) not justified</td>
<td>(a) controlled for aphasia*</td>
<td>6/10</td>
</tr>
<tr>
<td></td>
<td></td>
<td>1. (a) comparability between respondents’ and non-respondents’ characteristics is established and satisfactory*</td>
<td>(a) validated measurement tool on follow up**</td>
<td>1. (c) self-report</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>2. (b) hospital control</td>
<td>(b) controlled for age, language*</td>
<td>2. Yes</td>
<td></td>
</tr>
<tr>
<td>Reference</td>
<td>Study Design</td>
<td>Selection of Participants</td>
<td>Justification and Appropriateness</td>
<td>Comparability</td>
<td>Screening Tool</td>
</tr>
<tr>
<td>----------------------------------------------------------------------------</td>
<td>--------------</td>
<td>-----------------------------</td>
<td>-----------------------------------</td>
<td>---------------</td>
<td>----------------</td>
</tr>
</tbody>
</table>
| Hilari & Northcott (2006)                                                  | III          | (b) somewhat representative of the average in the target population | (a) justified and appropriate | 1. (b) comparability between respondents and non-respondents is not satisfactory
2. no control | (a) screening tool with moderate validity | (a) controlled for aphasia | (b) controlled for post-stroke time period and pre-morbid conditions | 1. (c) self-report | 2. N/A | (a) clearly described and appropriate |
| Matos et al. (2014)                                                       | III          | (c) selected group of participants | (b) not justified | 1. (a) comparability between respondents’ and non-respondents’ characteristics is established and satisfactory
2. (c) No description | (a) validated measurement tool** | (a) controlled for aphasia | (b) controlled for age, language, living place, cognitive and hearing functioning | 1. (b) self-report | 2. N/A | N/A |
| Naess et al. (2009)                                                       | III          | (c) selected group of participants | (b) not justified | 1. (a) comparability between respondents’ and non-respondents’ characteristics is established and satisfactory
2. (b) hospital control | (b) non-validated assessment tool for aphasia, but the tool is available | (a) controlled for aphasia | (b) controlled for age* | 1. (c) self-report | 2. Yes | (a) clearly described and appropriate |
| Łapkiewicz & Grochmalbach (2008)                                          | III          | (c) selected group of participants | (b) not justified | 1. (c) no description of the response rate or the characteristics of the responders and non-responders
2. (b) hospital control | (a) validated measurement tool** | (a) controlled for aphasia | 1. (b) self-report | 2. Yes | (a) clearly described and appropriate |
| Lee et al. (2015)                                                         | III          | (c) selected group of participants | (b) not justified | 1. (c) no description of the response rate or the characteristics of the responders and non-responders
2. (b) hospital control | (a) screening tool with moderate validity | (a) controlled for aphasia | (b) controlled for language, time post-stroke, pre-morbid conditions, functioning | 1. (c) self-report | 2. Yes | (a) clearly described and appropriate |
<table>
<thead>
<tr>
<th>Study</th>
<th>Quality</th>
<th>Group Selection</th>
<th>Response Justification</th>
<th>Comparator</th>
<th>Measurement Tool</th>
<th>Control</th>
<th>Analysis/Method</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dalemans, de Witte, Wade et al. (2010)</td>
<td>III</td>
<td>(c) selected group of participants</td>
<td>(b) not justified</td>
<td>1. (b) comparability between respondents and non-respondents not satisfactory 2. (c) no control</td>
<td>(a) screening tool with moderate validity*</td>
<td>(a) controlled for aphasia* (b) controlled for time post-stroke, age, living place, caregiver*</td>
<td>1. (c) self-report 2. N/A</td>
<td>N/A No statistical analysis. Qualitative analysis with use of an online software program. Description fair.</td>
</tr>
<tr>
<td>Fotiadou et al. (2014)</td>
<td>III</td>
<td>(c) selected group of participants</td>
<td>(b) not justified</td>
<td>1. (a) comparability between respondents’ and non-respondents’ characteristics is established and satisfactory* 2. (c) no control</td>
<td>(c) no description of a measurement tool</td>
<td>(a) controlled for aphasia* (b) controlled for language, age, content*</td>
<td>1. (c) self-report 2. N/A</td>
<td>N/A No statistical analysis. Framework analysis completed. Method of analysis qualitative and well described</td>
</tr>
<tr>
<td>Pommerehn et al. (2016)</td>
<td>III</td>
<td>(c) selected group of participants</td>
<td>(b) not justified</td>
<td>1. (c) no description of the response rate or the characteristics of the responders and non-responders 2. (c) no control</td>
<td>(c) no description of the measurement tool</td>
<td>(a) controlled for aphasia* (b) controlled for age, participation in therapy*</td>
<td>1. (c) self-report 2. N/A</td>
<td>N/A No statistical analysis. Thematic analysis based on ICF.</td>
</tr>
</tbody>
</table>
As evident in Table 2, all the studies were rated as Level III according to the ASHA rating scale, which is classified as ‘well-designed, non-experimental studies’. On the NOS rating scale a wide range of levels of evidence were obtained, with the studies displayed from highest to lowest levels of evidence, and alphabetically where appropriate. For the purpose of interpretation, a score of 0-3/10 was classified as a low level of evidence, a score of 4-6/10 as a moderate level of evidence and 7-10/10 as a high level of evidence. Two studies obtained a high level of evidence, six a moderate level of evidence and three a low level of evidence.

2.7. Data analysis

Thematic analysis was used to organise and synthesise the information obtained from the five qualitative and six quantitative studies, in accordance with the social participation domains identified in the ICF framework, namely domestic life, interpersonal relationships, education and employment, as well as community, civic and social life. Thematic analysis is a method used to interpret both implicit and explicit data items, requiring interpretation of the findings, which in this research study requires interpreting and synthesising the data into domains of social participation (Guest, MacQueen, & Namey, 2012). The contextual factors impacting on social participation is an additional aspect identified in this review. This aspect was highlighted as an area for further research in the Dalemans et al. (2008) review and recent research has since been conducted on this topic.

Attention was given to the reliability and validity of the systematic review with the use of the latest guidelines and checklists. The PRISMA-P checklist provided detailed guidance on relevant information that must be included in the review, as well as the methods that should be followed to ensure a high degree of transparency in the study and allow replicability of the review (Shamseer et al., 2015). An adapted version of the NOS (Wells et al., 2014) was used to provide an objective representation of the quality of evidence of the selected studies. The article as submitted to ‘Topics in Stroke Rehabilitation’ is included in Chapter 3, with formatting and referencing as outlined by the journal editors. See Appendix C for proof of submission.
CHAPTER 3

Aim of the chapter:

To present the systematic review article as submitted to *Topics in Stroke Rehabilitation* on 11/05/2017.

3.1. Article
Social participation in working-age adults with aphasia: An updated systematic review

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Co-author: Bhavani Pillay; University of Pretoria, Private Bag X20, Pretoria, Hatfield, 0028; 012 420 4919; bhavani.pillay@up.ac.za

The research was planned by all three authors, conducted by Caitlin Pike and the article written up by Caitlin Pike, Alta Kritzinger and Bhavani Pillay.
Social participation in working-age adults with aphasia: An updated systematic review

Background: A previous systematic review found limited data regarding social participation in working-age people with aphasia (PWA). This population has many roles to fulfill, that are negatively affected by aphasia. A review of recent studies may reveal more information on challenges in re-establishing social roles and thus may inform treatment thereof.

Method: The aim was to provide an updated systematic review on social participation in PWA under 65 years of age. Studies from 2005-2017 were searched from Scopus, Pubmed and Psychinfo. Search terms were derived from the International Classification of Functioning, Disability and Health (ICF) and the Aphasia- Framework for Outcomes Measures (A-FROM). Aspects of domestic life, interpersonal relations and interactions, education and employment and community, civic and social life were investigated.

Results: From 2,864 initial hits, 11 studies were identified, all of which were on the American Speech-Language-Hearing Association (ASHA) Level III of evidence. The studies indicated that participation in domestic life is reduced and PWA showed reduced social networks, loss of friendships and changes in the quality of marital relations. Few PWA returned to work or spent time on education. Limitations in community, civic and social life were noted and there were contradictory findings on the impact of contextual factors on social participation. There was an increase in research into contextual factors impacting on social participation in PWA and in the use of conceptual frameworks in the last decade.

Conclusions: Social participation in working-age adults is limited across the social domains. While the ICF conceptual framework is increasingly used, no studies used the A-FROM. There is greater use of standardised assessments and larger sample sizes.

Keywords: stroke, aphasia; social participation, working-age, young, middle-age, ICF

Introduction

There is an increase in stroke incidence between the ages of 20 and 54\(^1\). This is of concern as individuals with stroke in young and middle adulthood face particular
challenges in regaining meaningful involvement in their roles as providers, partners and parents, which may no longer be required in later years\textsuperscript{2,3}. Aphasia is one of the strongest predictors of poor functional recovery following a stroke\textsuperscript{4} and impacts on many facets of daily life.

A number of participation intervention approaches have been developed, the goal of which are to maximize re-engagement in daily life\textsuperscript{5,6}. One approach, the Life Participation Approach to Aphasia [LPPA], emphasizes the concerns of the PWA within the treatment process\textsuperscript{7}. Another, the World Health Organization International Classification of Functioning, Disability and Health [WHO- ICF]\textsuperscript{7}, helps to re-direct the focus to health, well-being and quality of life (QOL)\textsuperscript{7}. An ICF compatible conceptual guide has been developed for aphasia: Living with Aphasia- Framework for Outcome Measurements [A-FROM]\textsuperscript{8}. This framework was developed to guide outcome measurements, with emphasis on real-life outcomes\textsuperscript{8}.

These approaches are reflective of goals identified by PWA and their families. A key area in which PWA would like to improve is life participation\textsuperscript{9,10} and this aspect is highlighted as fundamental to living successfully with aphasia\textsuperscript{11}. Of concern is the finding that SLTs continue to be anchored in traditional medical models of treatment\textsuperscript{12}.

Consistent with the development of and subsequent drive towards the use of these approaches, there appears to be increasing research into social participation or QOL of PWA\textsuperscript{2,13,14}, however there appears to be limitations in the literature. In their systematic review (1960-2005) on social participation of PWA under 65 years, Dalemans et al. (2008) found that restrictions are evident across the social participation domains. The authors identified the need for increased use of a clear conceptual framework and well-defined concepts\textsuperscript{2}. They found that many studies had small sample sizes and used few standardized assessments. The researchers were unable to identify
studies describing community, civic and social life, one of the domains in “Activity and Participation” of the ICF, and they note a lack of research into the impact of the environment on social participation and return to work. It is evident that much is still unknown about the participation of working-age PWA.

When considering the dearth in literature, the increasing incidence in as well as the significant impact of aphasia on working-age adults, an expanded understanding of the social participation of these individuals is needed. With this knowledge, assessment and treatment procedures can be adjusted to the unique needs of this population for improved functional outcomes. This leads to the research question: What is known about the impact of aphasia on social participation in working-age adults with stroke-related aphasia and what is the level of evidence of these studies from the last decade?

Method

Study design

A systematic review was completed according to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses Protocols [PRISMA-P] checklist.

Study inclusion criteria

Three electronic databases were searched, based on relevance to the subject field: Scopus, PsychINFO and PubMed. The main search terms were: “aphasia”, and/or “stroke”, together with terms related to social participation according to the ICF and A-FROM frameworks: domestic life, relationships, education, employment, leisure, community life, social life and civic life. Additional terms included “quality of life”, “long-term outcomes”, “well-being” and “self-esteem”.

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Study selection

Three searches were conducted with the last search being run on 24/01/2017. Using this electronic search strategy (limits: 2005-2017, English, original articles), the initial search yielded 2,864 articles. This time limit was set as the review completed by Dalemans et al. (2008) included studies until 2005. A software program, Covidence, was used to synthesize searches, identify duplicates and review articles. Following removal of duplicates, 1,625 articles remained. The articles’ titles and abstracts were reviewed and articles that discussed aphasia in conjunction with an identified social participation aspect were included for full-text review. Forty-one articles met the criteria (Figure 1).

The following exclusion criteria were identified for the final review phase:

- Fewer than six PWA of 18 and 65 years, to eliminate single case studies which represent the lowest level of evidence on the American Speech-Language Hearing Association (ASHA) evidence rating scale, and may not significantly contribute to a systematic review.
- Percentage of PWA was less than 10% in quantitative studies.
- Mean age of the PWA was more than 65.
- PWA were not separately outlined in the population characteristics.
- PWA were not stroke-related.
- Participants presenting with aphasia after more than one stroke.
- Only the acute phase (0-3 months) was reported on.
- The measurement instrument, methodology and/or statistical/qualitative analysis were not described.
- Spouses or carers described their own needs in relation to the PWA.
- QOL was measured, but findings were not described in terms of social participation
- Social participation aspects were not separately described in the study outcomes
- Social participation was discussed according to changes following a treatment program

**Data collection process and data items**

Data was extracted from the 11 selected articles and compiled into pre-developed tables. These tables were structured according to the ICF social participation domains, as initial article review indicated that this continues to be the predominant framework used, as opposed to the A-FROM. The data items investigated are as follows:

3. Characteristics of the study, including title, author(s), year of publication, country, type of study, sampling method, mean age, sample size, social participation domain investigated, measurements used and the assessment period post-stroke (Table 1)

4. The impact of aphasia on social participation domains: domestic life, interpersonal life, education and employment, community, civic and social life.

**Risk of bias in selected studies**

An adapted version of the Newcastle-Ottawa Quality Assessment Scale [NOS] was used (Appendix B). This tool was developed to assess the quality of non-randomised studies. Each study was judged on three broad categories: study group selection, group comparability and outcomes with a greater number of stars indicating a higher level of evidence. The content validity and inter-rater reliability of this rating scale have been established.
The widely accepted American Speech-Language-Hearing Association [ASHA] level of evidence rating scale were also used to categorise the selected studies (ASHA, 2004). This scale rates studies on four levels based on the research design used, with the lowest rating being IV and the highest being I (see Table 2). The ratings were independently completed by two of the researchers, and differences noted were resolved after discussion.

As evident in Table 2, all the studies were rated as Level III according to the ASHA rating scale, which is classified as ‘well-designed, non-experimental studies’. On the NOS rating scale a wide range of levels of evidence were obtained, with the studies displayed from highest to lowest levels of evidence, and alphabetically, where appropriate. For the purpose of interpretation, a score of 0-3/10 stars was classified as a low level of evidence, a score of 4-6/10 stars as a moderate level of evidence and 7-10/10 stars as a high level of evidence. Two studies obtained a high level of evidence, six a moderate level of evidence and three a low level of evidence.

Data analysis

Thematic analysis was used to organize the data. This was done according to the social participation domains identified in the ICF. In addition, the contextual factors impacting on social participation were investigated.

Results

Domestic life

Domestic activities include social activities (e.g. shopping, home-making) and other secondary activities of daily living. Studies that met the criteria were published from 2010 to 2016 (Table 3). The Echelle de Communication Verbale de Bordeaux (ECVB) was used in two studies that were part of a single cohort study and that were on
moderate and high levels of evidence \textsuperscript{18,19}. This questionnaire investigates communication activity in daily living. The factors identified as being the most challenging for working-age PWA in both studies were: using the phone, using checks and credit cards and communicating in social activities. Activity limitations in communication were more severe for individuals with severe aphasia than those with moderate aphasia\textsuperscript{19}. Two studies\textsuperscript{20,21} used the Community Integration Questionnaire (CIQ)\textsuperscript{22,23}, a tool divided into three subscales: Home Integration, Productivity and Social Integration. The Home Integration subscale indicates the frequency of participating in activities such as shopping and housekeeping. Both studies, one on a moderate level and the other on a low level of evidence, found low scores in home integration. The final studies with low and moderate levels of evidence used a semi-structured interview\textsuperscript{24}, and the ICF checklist\textsuperscript{25}. These studies found limitations in domestic tasks, with Pommerehn et al. (2016) identifying predominant difficulties in meal preparation, household chores, goods and services acquisition and helping others.

**Interpersonal relations and interactions**

This domain includes all formal and informal relationships\textsuperscript{2}. This was investigated in six studies from 2006 to 2016 (Table 4). Two studies conducted analyses according to the ICF\textsuperscript{24,25}. One used a social network questionnaire and the Medical Outcome Study Social Support Survey (MOS-SSS)\textsuperscript{26}. The Dyadic Adjustment Scale (DAS) and Marital Communication Questionnaire (MCQ) were used in one study\textsuperscript{27} and another used the Communication Effectiveness Index (CETI)\textsuperscript{28}. The final study used a framework analysis of blog content\textsuperscript{29}.

On a moderate level of evidence, significant differences between PWA and people with stroke, and no aphasia were found with regard to social isolation (p=0.054; Fisher exact)\textsuperscript{28}. Aphasia impacts on numerous relationships. In marriage, PWA identify
a loss of harmony, satisfaction, cohesion, emotional expression and a loss in faith in the value of marriage\textsuperscript{27}. The marital changes appear to be more extensive for PWA than for individuals with stroke, but no aphasia\textsuperscript{27}. Interestingly, 86.3\% of PWA found their relationships with their children were maintained or improved following the stroke, and 75.4\% of PWA reported this to be true for relations with other relatives\textsuperscript{26}. A number of the identified studies, on a moderate to low level of evidence, indicated a reduction in social networks of PWA, particularly friendships\textsuperscript{24–26,29}. Studies with a low level of evidence note limitations in all relations\textsuperscript{25} and found particular challenges with in-depth conversations and participation in family activities\textsuperscript{29}. PWA experienced a higher degree of dependence and changes in the roles they played\textsuperscript{29}.

**Education and employment**

Education includes informal, vocational training and higher education, while employment consists of informal, remunerative and non-remunerative employment, excluding domestic work\textsuperscript{2}. These seven studies (2006-2016) are reflected in Table 5. Two studies on a moderate to high level of evidence used the Productivity subscale of the CIQ, with both finding limitations in productivity\textsuperscript{20,21}. Attention was given to return to work\textsuperscript{13,24,28,29} and the ability to participate in work activities\textsuperscript{24,25}. Overall, the studies indicated a greatly reduced productivity level, with few working-age participants returning to paid employment. In a study on a moderate level of evidence, it was found that PWA spent less time on education than individuals without aphasia\textsuperscript{20}. The data from an additional study with a low level of evidence indicated that five out of twelve PWA noted moderate or severe limitations in informal education\textsuperscript{25}.

**Community, civic and social life**

This includes leisure and recreational activities\textsuperscript{2}. The four studies (2010-2016) are
shown in Table 6. In two studies with moderate and high levels of evidence respectively, the Social Integration subscale of the CIQ was used\textsuperscript{20,21}, which indicated reduced social integration in working-age PWA, with a decrease in time spent on leisure activities\textsuperscript{20}. The remaining two studies on moderate and low levels of evidence respectively, used the ICF to report on their findings\textsuperscript{24,29}. Of the aspects investigated, restrictions in recreation and leisure activities were noted. Restrictions in community life, political life and citizenship were also found\textsuperscript{25}.

**Contextual factors**

Three studies (2010-2016) investigated contextual factors impacting on social participation of PWA\textsuperscript{21,25,30} (Table 7).

A number of personal factors were identified to impact on participation in young PWA: motivation, physical and psychological condition and communication skills\textsuperscript{30}. Environmental factors found to impact the PWA were the role of the central caregiver, characteristics of the communication partners (willingness, skills, knowledge) and quietness and familiarity of the living place\textsuperscript{30}. The support of individuals with whom the PWA lives had a positive impact on social participation\textsuperscript{30}. Furthermore, PWA found additional barriers to social participation to be services, systems and re-integration policies in employment\textsuperscript{25}. These personal and environmental factors were identified in two low level of evidence studies. Contrastingly, in a study with a high evidence rating, contextual factors were not significantly associated with social participation in aphasia\textsuperscript{21}.

**Discussion**

**Main findings**

A systematic review was completed to investigate social participation in four life
domains (domestic activities, interpersonal relations and interactions, education and employment and community, civic and social life) in working-age adults (18-65 years) with aphasia. Eleven articles were identified to be pertinent to the topic.

There appears to be limitations in the knowledge regarding the social participation of this population. Overall, it is evident that social participation across the four domains is greatly reduced for young PWA.

Domestic life has been relatively well researched, with varied concepts investigated. Three assessment tools/frameworks were used, namely the ICF, CIQ and the ECVB. Of those studies using the ICF, one mapped the participants’ responses onto the framework, with findings focused largely on household tasks, while the other study included a checklist that investigated a number of concepts (e.g. acquisition of goods and services, preparing meals). The CIQ investigates concepts such as preparing meals, housework and caring for children and the ECVB investigates concepts such as conversing with family members, making phone calls and using a credit card, from a more communicative perspective. While certain items overlap, the concepts assessed are varied and are reflective of the ICF. It is evident that domestic life for young individuals is negatively impacted by aphasia. This finding is in agreement with those found by Dalemans et al. (2008) in their systematic review. Dalemans et al. (2008) found nine studies in this domain and the current study identified six, however a greater percentage of the studies in the current review made use of standardized assessments.

Interpersonal relations and interactions in PWA are affected, with PWA reporting a reduction in the number of people within their social network, with particular loss of friendships, and negative changes in the quality of their marital relationships. Two of the six studies were analyzed according to the ICF, with consensus across the studies regarding the negative impact of aphasia on this life
domain. Although a wide variety of concepts were investigated, the findings are reflective of those found by Dalemans et al. (2008).

A number of studies investigated education and employment, with two studies using the ICF. Two studies used the Productivity subscale of the CIQ, which investigated work, training and volunteer programs and is well-reflective of the ICF. The remaining studies used interview formats or patient history reviews. Few PWA returned to work and many reported changes in their ability to participate in work tasks/productive activity. Two studies made mention of education, reporting that PWA spent less time on education than individuals without stroke and that some PWA felt moderate to severe limitations in their ability to participate in educational-type activities. The attention to education appears to be a new avenue of research in PWA, as Dalemans et al. (2008) did not identify findings on this topic in their review.

The domain with the most limited research was that of community, civic and social life. Two studies reported on the Social Integration domain of the CIQ, which investigates items such as going out and leisure activities. Both studies found limitations in social integration, which was also noted by a study analyzing interview findings of participants according to the ICF. Dalemans et al. (2008) found similar results in leisure activities, but were unable to identify research on the remaining aspects of this domain. In the current review, only one study commented on all aspects of this domain (community life, recreation and leisure, religion and spirituality, political life and citizenship). It appears that PWA found greatest limitations in community life, followed by recreation and leisure and political life and citizenship. Few individuals found limitations in religion and spirituality.

An area of more recent research is the impact of contextual factors on social participation in PWA. While one study did not find that contextual factors impact on
social participation, the remaining two studies did, with a key environmental factor in both studies being the support of the central caregiver of the PWA.

The current review is in agreement with many of the findings reported in Dalemans et al. (2008). Of interest is the attention given to items identified as limitations or areas for further research. The researchers of the previous review indicated the need for increased use of a conceptual framework. It is evident that the ICF is becoming more widely used, which may enable more comparable results, however, the use is still fairly limited. It was noted that the A-FROM is not being used in research in this population. As this framework was developed in 2008, it would be expected that it would be more frequently used in recent years. Dalemans et al. (2008) also identified the need for research into the environmental factors impacting on social participation, and although limited, there is increasing research into this aspect. Two studies included comparison of social participation in stroke PWA, to stroke participants without aphasia, which was identified as a component for further research. While a number of self-developed questionnaires were used in the studies included in this review, there appears to be an increased trend towards using more standardized assessment measures. An encouraging factor is the apparent larger sample sizes in the included studies, which contributes to the reliability and generalizability of the findings. It appears that this rising population of young PWA is receiving increasing attention.

**Study limitations**

While the use of a conceptual framework is more evident than in the Dalemans et al. (2008) review, there were a number of studies in which the concept of participation was not well defined, which limited data synthesis. There still appears to be limited research focusing exclusively on working-age PWA. While a mean age limit of 65 was set, and the younger population is highlighted with this criterion, it is not possible to set a clear
distinction between the young and old population of PWA as yet. While every effort has been made to identify all studies related to social participation, limitations in search terms and the databases selected may have resulted in pertinent studies not being included.

**Future research**

While there is increased use of the ICF in studies, additional research is warranted to allow a more in-depth understanding of social participation, with a clear conceptual framework (ICF or A-FROM). The research on the impact of contextual factors on social participation appears limited and contradictory. All the studies included in the review were conducted in high-income countries, with the exception of one conducted in Brazil, which is an upper-middle income country\textsuperscript{31}. Social participation is likely to be influenced by contextual factors, such as a county’s per capita income. It is clear that studies from low to middle income countries is largely missing from existing data. While a few studies compare the social participation of young PWA to young adults with stroke, and no aphasia, this requires additional investigation.

**Conclusion**

Working-age PWA experience limitations across social participation domains. There is contradictory information regarding the impact of contextual on social participation in PWA, although a number of factors have been identified. The ICF is being used more frequently to guide research studies in social participation, but there is still a lack of consistency in the use of concepts, and to a certain extent, standardized assessment tools. These factors lead one to interpret the findings with caution.
Geolocation information

This systematic review was conducted in Pretoria, South Africa.

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Competing interests

The authors declare that they have no financial or personal relationships that may have inappropriately influenced them in the writing of the article.

Authors’ contributions

C.P. (University of Pretoria) main researcher. C.P. (University of Pretoria), A.K. (University of Pretoria) and B.P. (University of Pretoria) planned the research and wrote the article.

References


<table>
<thead>
<tr>
<th>Author and year of publication</th>
<th>Social participation aspect/domain(s) studied</th>
<th>Q/QL</th>
<th>Country</th>
<th>Sampling method</th>
<th>Mean age in months (R/SD)</th>
<th>Sample size</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dalemans, De Witte, Wade, &amp; Van den Heuvel (2010)</td>
<td>Contextual factors impacting on social participation</td>
<td>QL</td>
<td>Netherlands</td>
<td>Purposive sampling, with criteria to include different aphasia severities, genders, levels of mobility and time post-onset</td>
<td>57.4 (R 45-71)</td>
<td>13 PWA, 12 Caregivers</td>
</tr>
<tr>
<td>Dalemans, De Witte, Beurskens, Van den Heuvel, &amp; Wade (2010)</td>
<td>Domestic life, Education and employment, Factors impacting on social participation</td>
<td>Q</td>
<td>Netherlands</td>
<td>Purposive sampling</td>
<td>64.2 (R 35-87)</td>
<td>150 PWA</td>
</tr>
<tr>
<td>Darrigrand et al. (2011)</td>
<td>Domestic life, Factors impacting on social participation</td>
<td>Q</td>
<td>France</td>
<td>Convenience sampling, part of a larger study of stroke patients with aphasia</td>
<td>Severe aphasia: 63.7 (SD 15.1) Moderate aphasia: 64.1 (SD 10.4)</td>
<td>27 PWA (severe), 9 PWA (moderate)</td>
</tr>
<tr>
<td>Fotiadou, Northcott, Chatzidaki, &amp; Hilari (2014)</td>
<td>Interpersonal relations and interactions, Education and employment, Community, civic and social life, Factors impacting on social participation</td>
<td>QL</td>
<td>Greece</td>
<td>Purposive sampling of blogs sustained by a sole author who had aphasia following a stroke, and which reflected on their social network</td>
<td>48.8 (R 26-69)</td>
<td>10 PWA</td>
</tr>
<tr>
<td>Hilari &amp; Northcott (2006)</td>
<td>Interpersonal relations and interactions, Education and employment</td>
<td>Q</td>
<td>United Kingdom</td>
<td>Cluster sampling framework</td>
<td>61.6 (R 21-92)</td>
<td>83 PWA</td>
</tr>
<tr>
<td>Lapkiewicz &amp; Grochmal-bach (2008)</td>
<td>Interpersonal relations and interactions</td>
<td>QL</td>
<td>Poland</td>
<td>Purposive sampling of married couples, with one partner with severe aphasia</td>
<td>58.7</td>
<td>22 PWA (and partner), 21 Stroke, no aphasia (and partner)</td>
</tr>
<tr>
<td>Lee, Lee, Choi, &amp; Pyun, (2015)</td>
<td>Domestic activities, Education and employment, Community,</td>
<td>Q</td>
<td>South Korea</td>
<td>Convenience sampling of patients with post-stroke aphasia (≥6 months)</td>
<td>59.2 (SD 7.2)</td>
<td>32 PWA, 42 Control, no,</td>
</tr>
<tr>
<td>Study</td>
<td>Topics</td>
<td>Methodology</td>
<td>Country</td>
<td>Sample Size</td>
<td>Control Group</td>
<td></td>
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</tr>
<tr>
<td>Matos, Jesus, &amp; Cruice (2014)</td>
<td>Domestic life, Interpersonal relations and interactions, Education and employment, Community, civic and social life</td>
<td>QL</td>
<td>Portugal</td>
<td>Unclear</td>
<td>65 (R 41-80)</td>
<td>14 PWA, 14 Family members, 10 SLT</td>
</tr>
<tr>
<td>Mazaux, Lagadec, De Sèze, Zongo, Asselineau, Douce, Trias, Delair, Darrigrand (2013)</td>
<td>Domestic life</td>
<td>Q</td>
<td>France</td>
<td>Convenience sampling, part of a larger study of stroke patients with aphasia</td>
<td>65.1 (R 13.5)</td>
<td>100 PWA</td>
</tr>
<tr>
<td>Naess, Hammersvik, &amp; Skeie (2009)</td>
<td>Interpersonal relations and interactions, Education and employment</td>
<td>Q</td>
<td>Norway</td>
<td>Purposive sampling from computer-based hospital registries</td>
<td>42 (R 24-49)</td>
<td>20 PWA, 175 Control with stroke, no aphasia</td>
</tr>
<tr>
<td>Pommerehn, Delboni, &amp; Fedosse (2016)</td>
<td>Domestic activities, Interpersonal relations and interactions, Community, civic and social life</td>
<td>QL</td>
<td>Brazil</td>
<td>Convenience sampling from members of a group</td>
<td>48.25 (R 25-67)</td>
<td>12 PWA</td>
</tr>
</tbody>
</table>

Abbreviations: CVA: Cerebral Vascular Accident, NPO: Non-Profit Organisation, OT: Occupational Therapist, PWA: Person with Aphasia, Q: Quantitative, QL: Qualitative, R: Range, SD: Standard deviation, SLT: Speech-language therapist
Table 2. Quality assessment and level of evidence of selected studies

<table>
<thead>
<tr>
<th>Source</th>
<th>ASHA level of evidence</th>
<th>Selection (Maximum 5 stars)</th>
<th>Comparability (Maximum 2 stars)</th>
<th>Outcome (Maximum 3 stars)</th>
<th>Total stars</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mazaux et al. (2013)</td>
<td>III</td>
<td>(b) somewhat representative of the average in the target population*</td>
<td>3. (a) comparability between respondents’ and non-respondents’ characteristics is established and satisfactory* 4. Controls</td>
<td>(a) validated measurement tool on follow up**</td>
<td>1. (c) self-report 2. N/A</td>
</tr>
<tr>
<td>Dalemans, de Witte, Beurskens et al. (2010)</td>
<td>III</td>
<td>(b) somewhat representative of the average in the target population*</td>
<td>(a) justified and appropriate* 1. (a) comparability between respondents’ and non-respondents’ characteristics is established and satisfactory* 2. (b) hospital controls</td>
<td>(a) controlled for aphasia* (b) controlled for age, language*</td>
<td>1. (c) self-report 2. Yes</td>
</tr>
<tr>
<td>Darrigrand et al. (2011)</td>
<td>III</td>
<td>(c) selected group of participants</td>
<td>1. (a) comparability between (a) validated measurement tool on follow up</td>
<td>(a) controlled for aphasia* (b) controlled for age, time post stroke and premorbid conditions*</td>
<td>1. (c) self-report 2. Yes</td>
</tr>
<tr>
<td>Study</td>
<td>Methodology</td>
<td>Justification</td>
<td>Comparability</td>
<td>Tool Description</td>
<td>Control Variables</td>
</tr>
<tr>
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<tr>
<td>Hilari &amp; Northcott (2006)</td>
<td>III</td>
<td>(b) somewhat representative of the average in the target population*</td>
<td>(a) justified and appropriate*</td>
<td>1. (b) comparability between respondents and non-respondents is not satisfactory 2. no control</td>
<td>(a) screening tool with moderate validity* (b) controlled for post-stroke time period and pre-morbid conditions*</td>
</tr>
<tr>
<td>Matos et al. (2014)</td>
<td>III</td>
<td>(c) selected group of participants</td>
<td>(b) not justified</td>
<td>1. (a) comparability between respondents’ and non-respondents’ characteristics is established and satisfactory* 2. (c) No description</td>
<td>(a) validated measurement tool** (b) controlled for age, language, living place, cognitive and hearing functioning*</td>
</tr>
<tr>
<td>Naess et al. (2009)</td>
<td>III</td>
<td>(c) selected group of participants</td>
<td>(b) not justified</td>
<td>1. (a) comparability between respondents’ and non-respondents’ characteristics is established and satisfactory* 2. (b) hospital control</td>
<td>(b) non-validated assessment tool for aphasia, but the tool is available* (a) controlled for age* (b) controlled for age*</td>
</tr>
<tr>
<td>Łapkiewicz &amp; Grochmal-bach</td>
<td>III</td>
<td>(c) selected group of</td>
<td>(b) not justified</td>
<td>1. (c) no description of the</td>
<td>(a) validated measurement (a) controlled for aphasia*</td>
</tr>
<tr>
<td>(2008)</td>
<td></td>
<td>participants</td>
<td>response rate or the characteristics of the responders and non-responders 2. (b) hospital control</td>
<td>tool**</td>
<td></td>
</tr>
<tr>
<td>Lee et al. (2015)</td>
<td>III</td>
<td>(c) selected group of participants</td>
<td>(b) not justified 1. (c) no description of the response rate or the characteristics of the responders and non-responders 2. (b) hospital control</td>
<td>(a) screening tool with moderate validity* (b) controlled for language, time post-stroke, premorbid conditions, functioning at assessment*</td>
<td>1. (c) self-report 2. Yes</td>
</tr>
<tr>
<td>Dalemans, de Witte, Wade et al. (2010)</td>
<td>III</td>
<td>(c) selected group of participants</td>
<td>(b) not justified 1. (b) comparability between respondents and non-respondents not satisfactory 2. (c) no control</td>
<td>(a) screening tool with moderate validity* (b) controlled for time post-stroke, age, living place, caregiver*</td>
<td>1. (c) self-report 2. N/A</td>
</tr>
<tr>
<td>Fotiadou et al. (2014)</td>
<td>III</td>
<td>(c) selected group of participants</td>
<td>(b) not justified 1. (a) comparability between respondents’ and non-respondents’ characteristics is established and satisfactory* 2. (c) no control</td>
<td>(c) no description of a measurement tool (a) controlled for language, age, content*</td>
<td>1. (c) self-report 2. N/A</td>
</tr>
<tr>
<td>Pommeren et al. (2016)</td>
<td>III</td>
<td>(c) selected group of participants</td>
<td>(b) not justified</td>
<td>1. (c) no description of the response rate or the characteristics of the responders and non-responders 2. (c) no control</td>
<td>(c) no description of the measurement tool</td>
</tr>
<tr>
<td>Author</td>
<td>Measurements instruments</td>
<td>Mean months post onset (R/SD)</td>
<td>Concepts</td>
<td>Results</td>
<td></td>
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<tr>
<td>-----------------------------</td>
<td>-------------------------------------------------------------------------------------------</td>
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<td>--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------</td>
<td></td>
</tr>
<tr>
<td>Mazaux et al. (2013)</td>
<td>Orgogozo’s score, Barthel Index, ASRS, BDAE (French version), TLC, ECVB</td>
<td>(R 12-18)</td>
<td>domestic activities</td>
<td>Most impaired activities: conversation on complex themes, using the phone for a meeting, using checks and credit cards, communicating during social activities. Least impaired: asking for daily living needs, talking about one’s wishes and purposes, expressing feelings, conversation with relatives, answering on a phone, reading time and reading family post/mail. ECVB scores associated with work status and type of job at inclusion, stroke severity at inclusion and follow-up, aphasia severity at inclusion and follow-up, auditory comprehension impairment on inclusion, BDAE items of auditory comprehension, fluency, naming, reading and writing at follow-up, mean number of SLT sessions, depression at follow-up.</td>
<td></td>
</tr>
<tr>
<td>Dalemans, De Witte, Beurskens et al. (2010)</td>
<td>Structured interview, FAST, Barthel Index, COOP-WONCA, Personal Factors Questionnaire, Environmental Factors Questionnaire, CIQ</td>
<td>90.6 (R 6-372)</td>
<td>home integration</td>
<td>Home integration score of 4.8 (max score: 12) on CIQ subscale (SD: 3.6 Range: 0-12).</td>
<td></td>
</tr>
<tr>
<td>Darrigrand et al. (2011)</td>
<td>Orgogozo’s score, Barthel Index, ASRS, BDAE (French version), TLC, ECVB</td>
<td>(R 12-18)</td>
<td>communication activities of daily life</td>
<td>Persons with severe aphasia present with severe activity limitations in communication, with performance three times lower than individuals with moderate aphasia and four times lower than individuals without aphasia. Aphasia severity and communication disability, but not non-verbal communication, at follow-up, are related to the initial severity of aphasia. Most impaired factors: using a phone, credit card, chequebook, reading and filling in administrative documents, communication behaviours in social life. Non-verbal communication performance was not associated with aphasia severity.</td>
<td></td>
</tr>
<tr>
<td>Matos et al. (2014)</td>
<td>In-depth semi-structured interview</td>
<td>29 (R 3-89)</td>
<td>domestic activities</td>
<td>PWA reported difficulties participating in household tasks, but this was not identified as the domain most impacted by the aphasia.</td>
<td></td>
</tr>
<tr>
<td>Lee et al. (2015)</td>
<td>Modified Barthel Index; FAST;</td>
<td>29.1</td>
<td>home integration</td>
<td>Home integration scores: PWA- Mean: 2.6, SD: 3.0, Control- Mean: 5.6,</td>
<td></td>
</tr>
</tbody>
</table>
| GDS, CIQ: SAQOL-39 (SD 20.6) | SD: 3.0. PWA spent less time on activities of market, finance, shopping. No significant difference observed in meal preparation and household activities. Factors impacting on home integration: activities of daily living and mobility.

Pommerehn et al. (2016) | ICF checklist 52 (R 26.4-136.8) | Predominant difficulties with meal preparation, household chores, goods and services acquisition and helping others.

Table 4. Interpersonal relations and interactions

<table>
<thead>
<tr>
<th>Author</th>
<th>Measurements instruments</th>
<th>Mean months post onset (R/SD)</th>
<th>Concepts</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hilari &amp; Northcott (2006)</td>
<td>SAQOL-39, MOS-SSS, a social network questionnaire</td>
<td>42 (R 13-250)</td>
<td>social networks, perceived social support, HRQL</td>
<td>86.3% of PWA reported maintained or increased contact with their children, and 75.4% with their relatives, while 63.9% reported less contact with friends, with 30% being unable to name a single close friend. Decreased social network size (Mean: 10; Mode 4), with the general population having between 8-15 members in their social network. Size of social network in women, social companionship and informational support associated with HRQL. Overall PWA felt well supported, particularly in tangible support.</td>
</tr>
<tr>
<td>Matos et al. (2014)</td>
<td>In-depth semi-structured interview Analyses according to ICF framework</td>
<td>29 (R 3-89)</td>
<td>interpersonal relationships</td>
<td>PWA reported significant changes in their relationships, and noted changes or loss of friendships.</td>
</tr>
<tr>
<td>Naess et al. (2009)</td>
<td>MASRS, Nottingham Health Profile questionnaire part I, SSS, NGA, CETI</td>
<td>≥ 60</td>
<td>social isolation</td>
<td>Aphasia is associated with social isolation (p= 0.054; Fisher exact).</td>
</tr>
<tr>
<td>Łapkiewicz &amp; Grochmal-bach (2008)</td>
<td>MMSE, GDS, CNBA, BNT (Polish version), TT, DAS, MCQ</td>
<td>6</td>
<td>marital coherence, perceived support, QOL</td>
<td>PWA found a loss of harmony, satisfaction, cohesion, emotional expression, faith in the value of marriage. PWA showed more extensive changes in the marriage as well as in more significant areas of marriage as compared to individuals with stroke and no aphasia.</td>
</tr>
<tr>
<td>Fotiadou et al. (2014)</td>
<td>Thematic analysis of blog content using Framework Analysis</td>
<td>≥ 12</td>
<td>social networks, social support, family dynamics; factors impacting on social relationships</td>
<td>In-depth conversations more challenging, reduced participation in family activities, higher degrees of dependence and changed family dynamics and roles. Contact with friends was reduced (communication and physical difficulties impacting). Wider social networks mostly reduced (related to reduced work and community activities as well as environmental barriers and fatigue). Other people’s reactions as well as support towards the PWA had a big impact on the individual.</td>
</tr>
<tr>
<td>Pommerenh et al.</td>
<td>ICF checklist</td>
<td>52</td>
<td>relationships</td>
<td>Predominant difficulties with informal and formal relations with family</td>
</tr>
</tbody>
</table>
and strangers.

Table 5. Education and employment

<table>
<thead>
<tr>
<th>Author</th>
<th>Measurements instruments</th>
<th>Mean months post onset (R/SD)</th>
<th>Concepts</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dalemans, de Witte, Beurskens et al. (2010)</td>
<td>Structured interview, FAST, Barthel Index, COOP-WONCA, Personal Factors Questionnaire, Environmental Factors Questionnaire, CIQ</td>
<td>90.6 (R 6-372)</td>
<td>productivity</td>
<td>Mean productivity score on subscale of CIQ (SD: 1.6, Range: 0-5) (max score=5).</td>
</tr>
<tr>
<td>Hilari &amp; Northcott (2006)</td>
<td>SAQOL-39, MOS-SSS, a social network questionnaire</td>
<td>42 (R 13-250)</td>
<td>return to work</td>
<td>No PWA returned to full-time employment. Only 6% of PWA were involved in part-time or voluntary work/students, and 56% were of working age.</td>
</tr>
<tr>
<td>Matos et al. (2014)</td>
<td>Semi-structured interview</td>
<td>29 (R 3-89)</td>
<td>employment</td>
<td>PWA reported loss of employment and changes in their ability to participate at work.</td>
</tr>
<tr>
<td>Naess et al. (2009)</td>
<td>MASRS, Nottingham Health Profile questionnaire part I, SSS, NGA, CETI</td>
<td>≥ 60</td>
<td>return to work</td>
<td>Of ischemic stroke patients employed before the incident, 33% of PWA and 69% of those without aphasia were employed on follow-up.</td>
</tr>
<tr>
<td>Lee et al. (2015)</td>
<td>Modified Barthel Index; FAST; GDS; CIQ; SAQOL-39</td>
<td>29.1 (SD 20.6)</td>
<td>productivity</td>
<td>Productivity scores: PWA (Mean: 0.3; SD: 0.8), Control (Mean: 2.8, SD: 1.8). PWA spent less time on education than the control group.</td>
</tr>
<tr>
<td>Fotiadou et al. (2014)</td>
<td>Thematic analysis of blog content using Framework Analysis</td>
<td>≥ 12</td>
<td>return to work, factors impacting return to work</td>
<td>2/10 individuals were employed post-stroke. Factors impacting return to work: aphasia, fatigue, older age, epilepsy, short attention span and difficulty multitasking.</td>
</tr>
<tr>
<td>Pommeren et al. (2016)</td>
<td>ICF checklist</td>
<td>52 (R 26.4-136.8)</td>
<td>employment</td>
<td>Predominant difficulties with basic economic transactions and performing paid work.</td>
</tr>
</tbody>
</table>

Table 6. Community, civic and social life

<table>
<thead>
<tr>
<th>Author</th>
<th>Measurements instruments</th>
<th>Mean months post onset (R/SD)</th>
<th>Concepts</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dalemans, de Witte, Beurskens, et al. (2010)</td>
<td>Structured interview, FAST, Barthel Index, COOP-WONCA, Personal Factors Questionnaire, Environmental Factors Questionnaire, CIQ</td>
<td>90.6 (R 6-372)</td>
<td>social integration</td>
<td>Mean social integration score of 8.4 (SD= 2.2, range: 2-12) (max score: 20).</td>
</tr>
<tr>
<td>Matos et al. (2014)</td>
<td>Semi-structured interview</td>
<td>29 (R 3-89)</td>
<td>recreation and leisure</td>
<td>PWA reported particular restrictions in participating in recreation and leisure activities.</td>
</tr>
<tr>
<td>Lee et al. (2015)</td>
<td>Modified Barthel Index; FAST; GDS; CIQ; SAQOL-39</td>
<td>29.1 (SD 20.6)</td>
<td>leisure activities, socialising</td>
<td>PWA spent less time on leisure activities than the control group. Social integration score: PWA (Mean 5.7, SD: 3.0), Control (Mean: 9.9, SD: 2.2). Social integration associated with QOL. Frequency of social contact with friends, number of places visited and attendance of meetings was decreased in PWA. Less going out that the control group.</td>
</tr>
<tr>
<td>Pommerehn et al. (2016)</td>
<td>ICF checklist</td>
<td>52 (R 26.4-136.8)</td>
<td>community life, recreation and leisure, religion and spirituality, political life and citizenship</td>
<td>Participants noted the greatest disability in community life, followed by recreation and leisure and political life and citizenship.</td>
</tr>
</tbody>
</table>

Abbreviations: Barthel Index: Barthel Activities of Daily Living Index, CIQ: Community Integration Questionnaire, FAST: Frenchay Aphasia Screening Test, GDS: Geriatric Depression Scale, R: Range, SAQOL-39: Stroke and Aphasia Quality of Life Scale-39, SD: Standard deviation
### Table 7. Contextual factors impacting on social participation

<table>
<thead>
<tr>
<th>Author</th>
<th>Measurements instruments</th>
<th>Mean months post onset (R/SD)</th>
<th>Concepts</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Dalemans, Witte, Beurskens, et al. (2010)</td>
<td>Structured interview, FAST, Barthel Index, COOP-WONCA, Personal Factors Questionnaire, Environmental Factors Questionnaire, CIQ</td>
<td>90.6 (R 6-372)</td>
<td>personal factors, environmental factors</td>
<td>Factors impacting social participation: age, gender, performance on fADLs and aphasia severity ($\beta=0.205$). Environmental and personal factors do not independently impact on social participation.</td>
</tr>
<tr>
<td>Dalemans, de Witte, Wade, et al. (2010)</td>
<td>FAST Pre-structured diary Semi-structured interview Focus group interview</td>
<td>61.5 (R 16-132)</td>
<td>engagement, personal factors, social factors, environmental factors,</td>
<td>The level of engagement in social activities is more important than the number of activities. Personal factors: motivation, physical and psychological condition, communication skills; Social factors: the role of the central caregiver, characteristics of the communication partners (willingness, skills and knowledge); Environmental factors: quietness and familiarity of living place.</td>
</tr>
<tr>
<td>Pommerenh et al. (2016)</td>
<td>ICF checklist</td>
<td>52 (R 26.4-136.8)</td>
<td>contextual factors</td>
<td>Most reported the following factors as facilitators of social participation: attitude, physical, practical or emotional support from people PWA lives with, however certain individuals noted these same factors as barriers to participation. Additional barriers identified: services, training policies, work and employment.</td>
</tr>
</tbody>
</table>

Abbreviations: Barthel Index: Barthel Activities of Daily Living Index, CIQ: Community Integration Questionnaire, COOP-WONCA: Darmouth Coop Functional Health Assessment Charts/Wonca, FAST: Frenchay Aphasia Screening Test R: Range, SD: Standard deviation
List of figures

Figure 1. PRISMA flow chart of selection process
Figure 1. PRISMA flow chart of selection process

PubMed, Scopus, PsychINFO
2005-2017, English, Original studies only
2,864 Citation(s)

1,625 Non-Duplicate Citations Screened

Inclusion/Exclusion Criteria Applied

1,584 Articles Excluded After Title/Abstract Screen

41 Articles Retrieved

Inclusion/Exclusion Criteria Applied

27 Articles Excluded After Full Text Screen

3 Articles Excluded During Data Extraction

11 Articles Included
CHAPTER 4

Aim of the chapter:

The aim of this chapter is to discuss the implications of the research findings, to describe limitations in the study, to identify future avenues of study and to draw a final conclusion.

4.1. Implications of the research findings

The use of social models, as opposed to medical models, for assessment and intervention is gaining momentum across many medical fields. A more holistic approach to managing clients is being embraced in order to encourage optimal recovery, participation and ultimately an improved QOL. In line with this change in approach, this systematic review was conducted to determine to what extent aphasia impacts on the social participation of working-age adults. Eleven studies meeting the criteria for this review on social participation were included and all were identified to be on a Level III ASHA evidence level, with variation on the NOS rating scale from a low score of 2/10 to a high score of 8/10, with six of the eleven studies obtaining more than 5/10. Four of the eleven studies used the ICF, or components thereof, to guide or structure the study or data interpretation (Darrigrand et al., 2011; Matos et al., 2014; Mazaux et al., 2013; Pommerehn et al., 2016), while no studies used the A-FROM. Dalemans et al. (2008) reported that only seven of the 18 studies included in their review were published after the development of the ICF, which limited the use of the framework, however all studies in the current review were published after this time. It appears that there continues to be limitations in knowledge regarding the social participation of this population of individuals. Overall, it is evident that social participation across the four domains is greatly reduced for working-age PWA.

Six studies investigating domestic life are in agreement that domestic life for young individuals is negatively impacted by aphasia in a variety of ways. Many of the factors investigated are required on a daily basis, including using a phone, filling in forms, shopping and communicating in social settings. Dalemans et al. (2008) also concluded that there is a decrease in domestic activities in this population. This
highlights the need to address functional communication skills in speech-language therapy intervention and to include the patient and family in goal setting. The PWA may identify simple daily skills that they wish to achieve and that may be targeted in treatment. As previously mentioned, research indicates that both PWA and their family members identify increased life participation as key areas in which they would like to improve (Isaksen, 2014). One cannot ignore the need to educate the client, families and public on the presenting condition in order to minimise limitations to participating in domestic life and to reduce barriers in the environment. Restrictions in resources of low income countries, such as South Africa, prove a greater challenge in facilitating this aspect.

Interpersonal relations and interactions in PWA are reduced, with a decline in the size of social networks, loss of friendships and changes in the quality of marital relations. This decline in social contacts was also found by Dalemans et al. (2008) in working-age PWA, however they also noted changes in the quality of interactions across friendships and family relationships, and not just marital relationships as the current study identified. Dalemans et al. (2008) found a decrease in sexual activity, an aspect that was not identified in the current review. This knowledge regarding the changes in interpersonal relations and interaction in PWA should lead one to focus on education of the public and family on aphasia and strategies to improve communication. This concept is frequently emphasised, but not always practiced (Roth & Worthington, 2005). Communication is the basis of relationships. In PWA, communication is altered, and without knowledge of how to facilitate communication, friends and family may easily become overwhelmed and avoid the challenging situations. Sufficient knowledge and skills will facilitate communication and therefore relationships with various people in the PWA’s life. Increased informational support and social companionships is associated with greater HRQL (Hilari & Northcott, 2006) and meaningful relationships to ‘living successfully with aphasia’ (Brown et al., 2012).

A number of studies investigated education and employment. Few PWA return to work and many report changes in their ability to participate in work tasks as well as productive activity, which are similar findings to what Dalemans et al. (2008) found. While Dalemans et al. (2008) did not find research regarding the changes in education in PWA, the current review found that PWA spend less time on education
than individuals without stroke and some PWA feel moderate to severe limitations in their ability to participate in educational-type activities, including informal education. These findings not only emphasise the need to address skills required for work or some form of productive activity in therapy, but also the need to educate society and make changes in the environment (Graham et al., 2011). The current review was investigating working-age adults. These individuals would most likely be employed or engaged in some form of productive activity, if they did not have aphasia, and yet there are severe limitations. Many of these individuals were the primary breadwinner in the family, and they may still be required to assist financially. Barriers in the workplace in addition to the individual’s symptoms of aphasia limit their ability to return to work or education. As indicated on the ICF, contextual factors have a great influence on participation (WHO, 2001). If one could reduce the contextual barriers, however, participation would be facilitated, and this concept is strongly emphasised by proponents of the social model of disability (Hallowell, 2017).

The domain with the most limited research in both the current review and Dalemans et al. (2008) was that of community, civic and social life. The studies indicated limitations in social integration, including aspects such as going-out and leisure activities, which is in accordance with the findings of Dalemans et al. (2008). While Dalemans et al. (2008) only found studies that commented on leisure activities, the current review found a slight increase in research in this domain. It appears that PWA found greatest limitations in community life, followed by recreation and leisure and political life and citizenship. Few individuals found limitations in religion and spirituality. While conclusions must be drawn cautiously due to the limited data, it appears that community, civic and social life is negatively affected in working-age PWA. This has implications for QOL and Cruice, Worrall, and Hickson (2006) recommend encouraging leisure activities in PWA. The same can be highlighted for the remaining aspects of community, civic and social life.

An area of more recent research is the impact of contextual factors on social participation on PWA. This was highlighted as an area for further research by Dalemans et al. (2008) as the researchers were unable to find studies investigating this topic. While one study did not find that environmental and personal factors impact on social participation, the remaining two studies did, with a key environmental factor facilitating participation being the support of the central
caregiver of the individuals with whom the PWA spends the most time. While research is limited, it is evident that one cannot ignore the barrier- or facilitating-effect contextual factors may have on the PWA (WHO, 2001). As discussed, the attention to contextual factors becomes particularly relevant in low income countries, where resources that may facilitate participation and integration are limited. Research into social participation in PWA in this context is important to determine the extent reduced resources may have on participation. The current review did not identify any studies conducted in Africa or South Africa on social participation in this population, which is of concern as the incidence of stroke in the young continues to rise in this context (Maredza et al., 2015).

The current review is in agreement with many of the findings reported in Dalemans et al. (2008). Of interest is the attention given to aspects identified as limitations or areas for further research in the Dalemans et al. (2008) review. The researchers of the previous review indicated the need for the increased use of a conceptual framework to aid defining of participation. While only four studies in the current review made use of the ICF, there appears to be a greater number of studies that made use of concepts of the ICF, which may enable more comparable results. The use of the IC is still relatively limited, however, and the A-FROM framework was not found to be used in any of the identified studies. Dalemans et al. (2008) also identified the need for research into the environmental factors impacting on social participation, and although limited, there is an expanding area of research into this aspect. Furthermore, two studies included comparisons of social participation in stroke PWA, as opposed to stroke participants without aphasia, which was identified as an area for further research (Dalemans et al., 2008). While a number of self-developed questionnaires were used in the studies included in this review, there appears to be an increased trend towards using more standardised assessment measures. An additional encouraging factor is the apparent larger sample sizes in the included studies, which contributes to the accuracy of the findings. It appears that this rising population of working-age individuals with aphasia is receiving greater attention.

It was noted in the current review, however, that the A-FROM, which is the conceptual framework developed by speech-language therapists specifically for aphasia in line with the ICF, does not currently appear to be used in research.
regarding social participation of working-age PWA. As this framework was developed in 2008, increased frequency of use would be expected, particularly as the framework specifically focuses on communication as a construct of social participation (Kagan et al., 2008). It may be that an increase in the use of frameworks for research and treatment would be facilitated with the A-FROM, as it is more tailored for aphasia than the ICF, which may previously have limited its use in previous research.

Overall, it appears that unanswered questions regarding participation in working-age adults are being investigated and new questions have arisen from the current updated review.

4.2. Limitations in the study

- Lack of consistency in the use of concepts:

While the use of a conceptual framework, namely the ICF, appears to be more evident, the use continues to be fairly limited and there are a number of studies in which the concept of participation, or the domains within participation, is not well defined. The use of the ICF serves to provide a clearer definition of concepts and therefore enables improved comparison between studies. A wide variety of assessment tools and questionnaires were used, which also contributes to variances in concepts used. These variations limit the ability to identify clear concepts in a number of studies and therefore to draw accurate conclusions.

- Few studies with populations limited solely to participants under the age of 65 years:

Only one study investigated participants under the age of 65 only, and only one study investigating individuals under the age of 68 only. The remaining studies all included a wide range of ages. While a limit of a mean age of 65 was set for the current study, and the younger population is highlighted with the use of this criterion, it was not possible to set a clear distinction between the young and old population of PWA at this stage.

- Studies included:
While every effort has been made to identify all studies related to social participation, limitations in search terms and the databases selected may have resulted in pertinent studies not being included.

4.3. Future research

Progress has been made in the understanding of social participation in working-age adults with aphasia over the last decade, however further research into this topic is recommended.

- Further research into the working-age population specifically, is warranted, with possible investigation into the unique differences in social participation as opposed to older PWA.
- Further research is warranted to allow a more in-depth and comprehensive understanding of social participation within the context of a standardised framework. To date, the ICF has been used, however, the A-FROM may facilitate this due to the adaptation of the framework from the ICF that makes it more appropriate for research in social participation in aphasia. This may encourage researchers to make use of a framework, as it may be more applicable than the ICF.
- From the current review, it appears that more attention has been given to the social participation domains of domestic life, interpersonal relations and employment in working-age adults than to education and community, civic and social life. These less investigated social domains are aspects that are important for the young population in ways that may no longer be required for older adults. A thorough understanding of the impact aphasia has on these domains in young adults will facilitate appropriate treatment thereof.
- Although there is emerging research into the impact of contextual factors on social participation, this is fairly limited and somewhat contradictory. Further research is required to identify the overall impact of these factors, as well as the specific factors that may impact on the PWA performance and progress.
- While a few studies exist that compare the social participation of young adult with aphasia to young adults with stroke, and no aphasia, this avenue of research is fairly limited. In order to understand the true impact of aphasia, one must compare PWA to those with stroke, but no aphasia.
• It would be useful to research how social participation may be influenced by the presenting sub-type of aphasia. There is a great difference between the presenting symptoms of fluent vs. no-fluent aphasias and these differences may be reflected in differences in social participation as well.

• A number of studies investigated the impact of different treatment approaches on social participation. It would be valuable to identify exactly which protocols have shown success in the aphasia population in order to facilitate participation.

• It is clear from this review that the research regarding social participation of young PWA is conducted primarily in middle to high income countries as compared to low income countries. Social participation and community reintegration is affected by personal, functional and environmental factors, which may be facilitated by resources that may be more readily available in middle to high income countries (Akosile et al., 2016). Due to the unique challenges in low and middle income countries, such as South Africa, further research is needed in these settings. A longitudinal study may be warranted due to the comparatively low incidence of stroke in the younger population as opposed to individuals over 65 years of age, as well as due to the challenges that a multi-lingual country poses to assessment and comparison of participants.

4.4. Conclusion

Working-age PWA experience limitations across the social participation domains, including domestic life, interpersonal relations and interactions, education and employment as well as community, civic and social life. There is contradictory information regarding the impact of environmental and personal factors on social participation in PWA, although a number of factors do appear to have been identified. The ICF appears to be used more frequently to guide research studies in social participation, but it is still fairly limited and there is still a lack of consistency in the use of concepts, and to a certain extent, use of standardised assessment measures, in a number of studies. These factors raise concerns which lead one to interpret the findings with some degree of caution.
REFERENCES


incidence, severity, fluency, etiology, and thrombolysis. *Stroke*, 37(6), 1379–1384. http://doi.org/10.1161/01.STR.0000221815.64093.8c


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meta-analysis protocols (PRISMA-P) 2015: elaboration and explanation. BMJ (Clinical Research Ed.), 349(3), g7647. http://doi.org/10.1136/bmj.g7647


LIST OF APPENDICES

Appendix A: PRISMA-P checklist
Appendix B: Newcastle-Ottawa Quality Assessment Scale adapted for cross-sectional studies
Appendix C: Proof of submission to Topics in Stroke Rehabilitation
## APPENDIX A: PRISMA-P checklist

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<th>Table 2</th>
<th>PRISMA-P (preferred reporting items for systematic review and meta-analysis protocols) 2015 checklist: recommended items to address in a systematic review protocol</th>
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<td>Update</td>
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<td>Contact</td>
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<td>Role of sponsor or funder</td>
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<td><strong>Introduction</strong></td>
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<tr>
<td>Rationale</td>
<td>6</td>
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<td>Objectives</td>
<td>7</td>
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<td>Selection process</td>
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<td>Data collection process</td>
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<td>Outcomes and prioritization</td>
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<td>Risk of bias in individual studies</td>
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<td>Data synthesis</td>
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<td></td>
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<td>Meta-bias(es)</td>
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<td>Confidence in cumulative evidence</td>
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</table>
APPENDIX B: Newcastle-Ottawa Quality Assessment Scale adapted for cross-sectional studies

Selection:
1) Representativeness of the sample:
   a) Truly representative of the average in the target population. * (all subjects or random sampling)
   b) Somewhat representative of the average in the target group. * (non-random sampling)
   c) Selected group of users.
   d) No description of the derivation of the included subjects.
2) Sample size:
   a) Justified and satisfactory. *
   b) Not justified.
3) Non-respondents:
   a) Comparability between respondents and non-respondents characteristics established and good response rate. *
   b) Not satisfactory response rate or comparability between respondents and non-respondents.
   c) No description of response rate or responders and non-responders characteristics.
4) Ascertainment of the exposure (risk factor):
   a) Validated tool. **
   b) Non-validated but available or described tool.*
   c) No description.

Comparability:
1) Comparability of subjects in different outcome groups on the basis of design or analysis. Confounding factors controlled.
   a) Study controls for the most important factor (select one). *
   b) Study control for any additional factor. *

Outcome:
1) Assessment of outcome:
   a) Independent blind assessment. **
   b) Record linkage. **
   c) Self report. *
   d) No description.
2) Statistical test:
   a) Statistical test used to analyze the data clearly described, appropriate and measures of association presented including confidence intervals and probability level (p value). *
   b) Statistical test not appropriate, not described or incomplete
APPENDIX C: Proof of submission to Topics in Stroke Rehabilitation

Submission Confirmation for Social participation in working-age adults with aphasia: An updated systematic review

Topics in Stroke Rehabilitation Journal Office <em>editorialmanager@up.ac.za>

11 May (2 days ago)

Dear Mr. Peng,

Your submission entitled "Social participation in working-age adults with aphasia: An updated systematic review" has been received by journal Topics in Stroke Rehabilitation.

You will be able to check on the progress of your paper by logging on to Editorial Manager as an author. The URL is <http://strokehab.edmgr.com/).

Your manuscript will be given a reference number once an Editor has been assigned.

Thank you for submitting your work to this journal.

Kind regards,

Topics in Stroke Rehabilitation