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1 Title

- 2 Facilitators and barriers to participation in mental wellbeing programs by older Australians
- 3 with vision impairment: community and stakeholder perspectives

4 Running title

5 Vision impairment and mental wellbeing programs

6 Authors

- 7 Lisa Dillon^{a b*} PhD, lisa.dillon1@unsw.edu.au
- 8 Diana Tang^c MND, diana.tang@sydney.edu.au
- 9 Gerald Liew^c PhD MBBS MMED, gerald.liew@sydney.edu.au
- 10 Maree Hackett^{b d} PhD, mhackett@georgeinstitute.org.au
- 11 Ashley Craige PhD, a.craig@sydney.edu.au
- 12 Bamini Gopinath^c PhD, bamini.gopinath@sydney.edu.au
- 13 Lisa Keay^{a b} PhD, I.keay@unsw.edu.au
- 14 a School of Optometry and Vision Science, Faculty of Science, UNSW Sydney, Postal:
- 15 School of Optometry and Vision Science, The University of New South Wales, Sydney
- 16 UNSW NSW 2052, Australia
- 17 b The George Institute for Global Health, Faculty of Medicine, UNSW Sydney, Postal: PO
- 18 Box M201 Missenden Rd NSW 2050, Australia
- 19 c Centre for Vision Research, Department of Ophthalmology, The Westmead Institute for
- 20 Medical Research, The University of Sydney, Sydney, Australia, Postal: Centre for Vision
- 21 Research, 176 Hawkesbury Road, Westmead NSW 2145
- 22 d Faculty of Health and Wellbeing, University of Central Lancashire, Preston, United
- 23 Kingdom, Postal: University of Central Lancashire, Preston, Lancashire PR1 2HE

- e John Walsh Centre for Rehabilitation Research, Northern Clinical School, Faculty of
- 25 Medicine and Health, The University of Sydney, Postal: Kolling Institute of Medical
- 26 Research, RNSH, St. Leonards, NSW, 2650
- 27 *corresponding author

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29 The authors declare no conflict of interest.

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Abstract

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Objective: Older adults with vision impairment experience high rates of mental health problems, but very few access psychological support. We investigated community and stakeholder perspectives of the barriers and facilitators to participation in mental wellbeing programs for older adults with vision impairment. Methods: Adults aged ≥50 years with vision impairment (community) were recruited from the client database, and low vision rehabilitation (LVR) professionals (stakeholders) from staff of a LVR provider. Participants completed one-on-one semi-structured interviews which were designed and analysed using behaviour change theory. Results: Twenty-nine participants were interviewed; 16 community members and 13 stakeholders. Both groups cited mental health problems as a major concern, with many stakeholders reporting the grief and distress associated with vision loss experienced by their clients as having a negative impact on their mental and physical health. Major barriers to participation in mental wellbeing programs included a lack of awareness and difficulties accessing such programs, with stakeholders adding that their clients' lack of insight into their own mental health problems may reduce motivation to participate. Facilitators to participation in programs included the appeal of social interaction and inspirational speakers. An appropriate intervention could overcome these barriers, or enhance participation through education, persuasion, incentivisation, modeling, environmental restructuring, training, and enablement. Conclusions: While barriers were discussed more than facilitators to participation, there was general support for mental wellbeing programs. This study provides guidance from stakeholders for the development of mental wellbeing programs to address mental health problems in the growing number of older adults with vision impairment.

Introduction

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Globally, it is estimated that the number of people living with vision impairment (encompassing low vision or blindness that cannot be corrected with glasses or surgery) will rise from 38.5 million in 2020 to 115 million people by 2050; most of which (78%) are aged 50 years or older [1]. Older age and vision impairment are associated with an increased risk of developing a mental health condition, particularly depression [2]. An estimated 69,519 non-Indigenous Australians aged 50 years or above and 4,282 Indigenous Australians aged 40 years or above were living with irreversible vision impairment in 2016 [3]; all at consequent risk of mental health problems. Older adults with vision impairment are approximately three times more likely to experience significant depressive symptoms, compared with those without vision impairment [2, 4]. This is attributed to the impacts of vision impairment on functional capacity and activities of daily living [5-8]. Research suggests that low vision rehabilitation (LVR) services, and psychological interventions involving self-management and problem-solving, may be effective in reducing depressive symptoms and depressive disorders among adults with vision impairment [9-12]. However, although 70% of adults with vision impairment report wanting psychological support, only 9% report receiving it [13], and it remains unclear why uptake of support is so low. One study of LVR professionals' (hereafter stakeholders) perspectives of a mental wellbeing program found a number of barriers to the problem-solving treatment for primary care (PST-PC) being delivered in a LVR setting, that could be overcome through professional training, support and improved screening [14]. The only other study reported on the reasons older adults with vision impairment withdrew early from PST-PC, and found responses ranging from the program being perceived as not relevant, to the program goals being achieved early [10]. Given this scant literature, the aim of this qualitative study is to investigate the barriers and facilitators to participation in a mental wellbeing programs, from the perspectives of

community-dwelling older adults with vision impairment (community members) and stakeholders.

Subjects and Methods

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This qualitative study involved semi-structured, one-on-one telephone or face-to-face interviews with community members and stakeholders, including orientation and mobility specialists and optometrists. This study was conducted within New South Wales (NSW) and the Australian Capital Territory (ACT), Australia. Community members and stakeholders were recruited using purposive sampling through invitation letters and follow-up phone calls between August and October 2019. To address the study aim, we expected to recruit up to 20 community members and up to 15 stakeholders for interview, or until data was saturated in each group, which was assessed by researchers (LD, LK) at regular meetings. Community members were recruited from the client database of an Australian vision rehabilitation organisation, Guide Dogs NSW/ACT. Clients that who had consented to be contacted regarding research were sent an email and invited to contact the research team directly to express interest in the study. Participants were required to be aged 50 years and older and speak conversational English. Stakeholder participants were employed in a client-facing capacity by Guide Dogs NSW/ACT, and were similarly sent an email and invited to contact the research team directly to express interest in the study. The behaviour change wheel [15] was used as a framework to design the interview guides (Supplementary File 1 and 2), and consequent analysis of the data. The behaviour of interest was participation in a mental wellbeing program. Specifically, the wheel uses the COM-B model to explain an individual's propensity to adopt a behaviour (see the inner wheel of Figure 1). Behaviour can be in turn targeted by nine intervention functions: coercion, education, enablement, environmental restructuring, incentivisation, modelling, persuasion,

restriction, training [15]. The wheel also includes policy categories which can impact the intervention functions; however, these were considered outside the scope of this study. Semi-structured interviews, lasting 15-45 minutes, were conducted by two Master of Clinical Optometry students (AW, RM) and a Faculty of Medicine and Health PhD student (DT). The students were given initial training and ongoing support by two experienced public health qualitative researchers (LD, LK). Interviews were audio recorded using digital recorders, transcribed verbatim, and analysed using NVivo software, using deductive analysis [16], following the COM-B and intervention functions of the behaviour change wheel [15]. We took an iterative approach to data analysis [17], whereby data were revisited, coded, and themes discussed many times to ensure analytic reflexivity. Transcripts were coded separately by two of the students (AW, RM). Initial coding was then presented and critiqued in a meeting including all three students, chaired by LD. Two of the students (AW, RM) then collaborated to establish themes under each subset of the COM-B model, which were discussed and agreed upon at regular fortnightly meetings with LD and LK. Themes were then coded using intervention functions, and categorised as barriers or facilitators to participation in a mental wellbeing program by LD and DT.

Patient and public involvement

No patient under medical care or members of the general public were involved in the design, recruitment or conduct of the study. However, there will be formal engagement with the LVR provider's advisory panel to discuss the next stages of developing and implementing a mental wellbeing program. Those participants who requested feedback will be informed via email or telephone call about the results.

Ethics approval

Ethics approval was granted by the University of New South Wales Human Research Ethics Committee (HC190356). <u>A participation information statement was sent to interested participants</u>, and oral informed consent was gained before commencement of the interview.

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The study is reported in line with the COREQ statement[18], supporting transparency in reporting of qualitative research.

Results

Of the 35 community members and stakeholders invited, 29 (83%, 16 community members and 13 stakeholders) completed an interview. The 13 stakeholders included 12 orientation and mobility specialists and one optometrist. All stakeholder interviews were conducted over the phone, while seven of the community participants requested face-to-face interviews in their homes due to difficulties with hearing and/or accessing a phone.

We identified twelve themes (Figure 1); eight represented barriers, and four facilitators to

mental wellbeing program participation. Illustrative quotes associated with each theme are presented below, with intervention functions in brackets next to the COM-B component. Participants are identified as C for community members and S for stakeholders, followed by

Both participant groups showed a high level of interest through long and engaging discussions around the mental wellbeing of people with vision impairment. Stakeholders expressed that many of their clients have concerns regarding their mental wellbeing, indicating the importance of this topic and need for such programs:

"I'm finding most of the clients that I...work with, express that they have anxiety, depression, or have had nervous breakdowns, or are currently having mental health issues." (S002)

Barriers

Physical capability (Enablement, Training)

156 Mobility impacts participation

an identification number.

Participants reported difficulties in mobility as a result of their vision loss which prevented them from engaging with their community and/or programs:

159	"Because you can't go out. That's the biggest problem" (C008)
160	"So, you don't tread on little kids or get tangled up in dog leadsthat's always in the
161	back of your mind. I think I'll just stay at home." (C004)
162	In particular, there were concerns regarding loss of the ability to drive:
163	"You're not going to get them in a group scenario, but also they can't drive so they can't
164	get to anywhere." (S008)
165	Some participants also found that their mobility limitations and location of residence
166	interfered with their ability to connect with other individuals with vision impairment and
167	contributed to feelings of isolation:
168	"I would love to have a group of other people who were also vision impaired, but they
169	seem to be scattered all over the country." (C020)
170	Vision loss impacts everyday activities
171	The inability to accomplish activities of daily living deterred participation in programs and
172	other activities:
173	"Because of their vision impairment, I don't think they integrate as much with others in
174	their community." (S009)
175	"I can't watch TV and I do like TV actually. I can't read anything anymore and I used to
176	love a newspaper." (C020)
177	Psychological capability (Education, Training, Enablement)
178	Ripple effects
179	Stakeholders expressed concerns about mental health problems extending into other
180	aspects of their client's lives and preventing participation in mental wellbeing programs:

181	"Stress and mental health, the physiological changes to the body impacts on people's
182	functional vision. There's all these ripple effects if the mental health component is not
183	addressed." (S006)
184	"I'd be the only person they'd see that week and maybe they'd have a cryneed to work
185	through the issues they have before they can start doing routes and getting out in their
186	community." (S015)
187	Self-perception of mental health
188	The acknowledgement of having a mental health problem varied among participants:
189	"One very good friend who's also legally blindwhen I bring up the subject with him he
190	says, Oh, better not even to think about it. And he's not really getting all the services
191	that he should be getting as a blind person." (C004)
192	"Has it affected me? I lost my licence. I'm pretty much housebound. My doctor wants
193	me to see a psychologist. And I said, No, I'm not that bad."" (C008)
194	"I try to prepare myself for the future, but I don't think it's affected my mental health, as
195	far as others are concerned anyway." (C009)
196	Physical opportunity (Environmental restructuring, Enablement)
197	Facilitation of services
198	Participants expressed concerns about their ability to access mental wellbeing programs
199	following their vision loss:
200	"When I lost my sight I had to really scrabble and call for information." (C007)
201	"If there is support, they [support provider] could call our clients[so] the client doesn't
202	have to initiate [seeking services] themselvesbecause I have a lot of clients who once
203	they lose their vision, they're not able to navigate a phone." (S003)

204	Stakeholders also expressed their concerns about now they can best assist their clients to
205	receive support for their mental health problems:
206	"If we had a script or a proper format that once a client has identified that they've got
207	mental health issues that we should say because you've mentioned that to us can we
208	write some information down and pass it on to someone to suggest help or at least to
209	refer." (S013)
210	Lack of awareness of services
211	Community and stakeholders were unaware of current mental wellbeing programs targeted
212	at Australians with vision impairment:
213	"I'm not aware of any program for people with vision impairment." (S012)
214	"In a major Sydney hospital and they didn't ever say to me that I needed to be referred
215	to Vision Australia or Guide Dogs or anywhere like that. And I just said, Okay, well, I can
216	do this. But the trouble is I couldn't see." (C005)
217	Social opportunity (Environmental restructuring, Enablement, Modelling)
218	Stigma about mental health problems
219	Participants described the negative stigma surrounding mental health issues as a barrier to
220	participation:
221	"it doesn't seem to be talked about a lot." (C015)
222	"People have a perception, there's still a stigma around mental illness, so a lot of them
223	might not disclose." (S005)
224	"I would say don't call it a mental health program." (C005)
225	Reflective motivation (Education, Persuasion, Incentivisation)
226	Grief associated with vision loss

227 Feelings of grief and beliefs regarding vision loss can make participation and engagement 228 unmotivating: 229 "If a person is depressed and feels like they can't do anything, then it's likely they're not 230 going to go into orientation or mobility training with high expectations, and a good 231 learning frame of mind." (S012) 232 **Facilitators** 233 Social opportunity (Environmental restructuring, Enablement, Modelling) 234 Community connections 235 The desire for a connection to the community particularly with those experiencing similar vision loss was described by participants: 236 237 "...having a group that comes together that has a focus on adapting to change and 238 understanding... I think is fairly powerful." (C015) "Although technology and online stuff is cool and innovative...it sort of takes away from 239 240 that human connection that you have when you sit in the same room as somebody." 241 (S015) 242 "There's others there to talk to and pass the time of day with. It's not a discussion group, 243 it's just to pass the time. You know, with the social aspect of it, but they're getting to 244 know me." (C003) 245 "For so long I just thought our family was a family of freaks because not knowing of 246 anyone else that had it...30 years ago I joined up with the Nepean Blind Sports Club...I 247 met a couple of people with the same condition and... went to social events out there." 248 (C012) 249 Who can facilitate mental wellbeing programs

250	Participants also expressed their opinions about who they think would be best suited to
251	deliver low vision mental wellbeing programs. Different levels of expertise were articulated
252	ranging from lay-facilitators to mental health professionals like psychologists:
253	"Anybody who has got group-based skills and some level of working with groups."
254	(C005)
255	"I always feel that if someone's been through a situation they're the ones I think are the
256	better ones." (C012)
257	A consultant psychologist with a good knowledge of grief it would be great to be able
258	to run scenarios by that person. And say, this is what I'm dealing with, or this is what I've
259	done. It would be lovely to be able to say to clients If you want to talk more about this,
260	we have a psychologist on staff that might be able to give you a ring." (S004)
261	"It would be good if we had one person that they could make a call to that has more
262	specific information and then can put them in touch with people in their own area or
263	what organisations are close to them." (S013)
264	Automatic motivation (Modelling, Enablement)
265	Inspiration/role model
266	Some participants stated how a role model is an inspiring motivation for participating in
267	mental wellbeing programs:
268	"Those have been some of our better meetings when we've been inspired by others in
269	other words." (C009)
270	"hearing people's stories about how they might have done it is really powerful for
271	people who might be going through that part of the process." (C005)
272	Reflective motivation (Education, Persuasion, Incentivisation)
273	Self-efficacy

Many participants believed they were self-sufficient and capable of handling their own issues:

"I can't change it, I've just got to learn to adjust to it and that's just going to be my life."

277 (C011)

"I don't feel sorry for myself, there's no point. I've got two legs, I can walk, so life's

good." (C022)

Discussion

Consistent with evidence of the high burden of mental health problems in older adults with vision impairment [2, 4], we found strong interest in developing mental wellbeing programs among community members and stakeholders. The interviews revealed that both groups acknowledged the substantial negative impacts of mental health problems. However, both groups were unaware of any mental wellbeing programs specific to people with vision impairment, thus, confirming the importance of developing a mental wellbeing program tailored to this population group.

We identified two times more barriers than facilitators (i.e. eight vs four) to effective participation in a mental wellbeing program. The intervention function *Enablement* was relevant to all three facilitators, and the majority of barriers. Enablement refers to increasing means, and reducing barriers [15], and is key to designing a program for this population. A major concern related to Enablement, along with Environmental Restructuring (changing the physical or social context [15]), was functional disability. Many community members felt that vision loss limited their capability and opportunity to access programs due to difficulty reading or finding information, and travelling to where programs were held. Travel concerns are a common barrier in this population [19], and is influenced by the severity of vision impairment, distance to program location, and availability of support people.

Recommendations to overcome these barriers, as they relate to intervention functions [15],

include: producing material in larger, easy-to-read print (*Enablement*); LVR professionals telling community members what options are available (*Education*; increasing knowledge or understanding); and program organisers making community transport available (*Environmental Restructuring*) or provide orientation and mobility services (*Training*; imparting skills), if travel is required.

In addition to physical limitations, the acknowledgement of having a mental health problem influenced participation and varied among participants. Some mentioning conditions like depression and anxiety, others downplaying any concerns and associating it with complaining, and the remainder stating that their vision loss did not have any impact on their mental health. The intervention functions of *Modelling* (an example for people to aspire to or imitate) and *Education* are particularly relevant to alleviate these concerns and key to tackling the barriers of social stigma and self-perception. Nyman et al. [20] identified that vision impairment can impact psychosocial well-being through social isolation; while acceptance of vision loss, and social support were facilitators of adjustment. We also identified self-efficacy as a facilitator, which has been shown to lead to good mental health outcomes in other contexts [21]. Given there was general consensus among both groups of the benefits of social connection and positive role models, it is recommended interventions include this in their design.

A pilot feasibility study in Australia recently investigated the delivery of PST-PC by LVR professionals to older adults with vision impairment who met the criteria for mild symptoms of depression [10]. PST-PC, based on Cognitive Behavioural Therapy (CBT) principles, is a low-intensity approach designed to assist with functional adjustment, resilience building and generalised well-being. PST-PC can be delivered by non-mental health professionals, face-to-face, as well as over the phone, or over video conference, significantly increasing its reach. Holloway et al. [10] found that those that who stayed in the study had significant reductions in depressive symptoms, and improvements in health-related quality-of-life and problem-focused coping. Those who withdrew from the study early were typically older, with

complex needs, as has been found in other studies [22-24]. Thus, it is critical that any future interventions must consider methods to retain adherence of older participants with competing health priorities.

Varying levels of mental health problems and requirements for support were reported by community members as well as stakeholders about their clients. A stepped-care approach may be the most efficient in this context, as not all clients require the same type or intensity of intervention [25]. Stepped-care comprises different intervention components, with the idea that if the first, less intensive step does not lead to a reduction in symptoms, then a person moves to the next step, consisting of more intensive and potentially more expensive interventions [9]. Several randomised controlled trials conducted outside the field of low vision have found that a stepped-care approach can be effective in minimising depression and/or anxiety [26, 27], and has been endorsed to address depression in older adults in clinical guidelines, such as the UK NICE guidelines [28]. The ideas for interventions identified in this study could be integrated within a stepped-care approach, including social groups, psychologist referral and PST-PC.

Strengths and limitations

This is the first study to explore the perspectives of older adults with vision impairment and client-facing professionals involved in LVR regarding mental health problems. The inclusion of client-facing professionals provides essential insight needed to design a stakeholder-driven intervention. A second strength is the semi-structured nature of the interviews which allowed for tailored discussions to explore each participant's personal outlook and interpretation of the impacts of mental health problems and vision impairment. However, this also resulted in interviews of varying length, and occasional tangents in conversation.

Despite compelling findings, the authors acknowledge that the study is limited to one community organisation and therefore the presented results are only a preliminary indicator of the mental health perspectives in this population group. Moreover, community member participants were recruited through purposive sampling from the client base of one

Australian LVR provider, and specifically only those clients who had agreed to be contacted about participating in research, and had also agreed to participate in this particular study. This potential selection bias may have influenced our results, and may limit the generalisability of findings to those not associated with an Australian LVR provider, or those not interested in participating in research or this particular study. Similarly, stakeholder participants were orientation and mobility specialists and one optometrist from the one Australian LVR organisation, which may limit the generalisability of findings to these LVR providers in other organisations, or other professions involved in LVR, such as occupational therapists, orthoptists, assistive technology specialists and social workers. Nonetheless, as data saturation was achieved, we are confident the range of themes and corresponding intervention functions suitably address the study aim. Moreover, as participants were recruited through purposive sampling, there may be influences of selection bias influencing results and perhaps likely underestimating the mental health problems of individuals not associated with an LVR provider. Furthermore, information on visual acuity, cause of vision loss and presence of comorbidities was not available for collection. This information may have been valuable in terms of interpreting the qualitative data. This study has several key implications for clinical practice. First, improving the mental wellbeing of visually impaired older adults is a priority. We postulate this may enhance their participation in other programs, such as orientation and mobility, and this in turn could improve their physical health, community connections, and overall well-being. Second, LVR professionals, such as orientation and mobility specialists, and optometrists, are in a unique position to identify those at risk of, or already experiencing mental health problems and can refer as necessary, if provided with the necessary training. Third, older adults with vision impairment may benefit from appropriate education and training to better support them as they adjust to living with a vision impairment. We suggest this may be in the form of evidence-based education regarding their vision diagnosis and prognosis, or psychological support from an appropriately trained mental health practitioner. Fourth, providing older

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adults with vision impairment the opportunity to connect with others in similar situations, as well as those who are living well with vision impairment, is likely to improve their community connections and social engagement. As a next step, the suggestions for future interventions collected here will be presented through round tables with older adults with vision impairment and service providers. This will ensure that any developed strategies are practical and acceptable to the community and stakeholders, with the aim to improve the ultimate adoption and scalability of a mental wellbeing program.

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Conflict of Interest

The authors declare that they have no conflict of interest.

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Titles and legends to figures

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Figure 1 Barriers and facilitators to mental wellbeing program participation mapped on the behaviour change wheel[15]. Barriers to participation appear in bold, while facilitators appear underlined, with their relative intervention function presented in the outer wheel.