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Perspectives of people with late age-related macular degeneration on mental health and mental wellbeing programs: a qualitative study

Journal:	<i>Ophthalmic and Physiological Optics</i>
Manuscript ID	OPO-OA-3032.R2
Manuscript Type:	Original Article
Date Submitted by the Author:	23-Nov-2020
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<p>Keywords:</p>	<p>Age-Related Macular Degeneration, mental health, qualitative study, wellbeing programs, depression</p>



Table 1 Demographic characteristics of participants (n=29)

Study ID	Age (years)	Sex	AMD type	Years since any AMD diagnosis	Visual Acuity	History of other eye conditions
001	83	Male	RE: late (geographic atrophy and neovascular) LE: late (neovascular)	RE: 3.5 LE: 4.5	RE: 6/60-3 LE: 6/36+2	Cataract in both eyes; penetrating keratoplasty in RE requiring corneal graft
002	79	Male	RE: late (neovascular) LE: late (neovascular)	RE: 16 LE: 12	RE: 6/60-3 LE: 6/60-1	Cataract in both eyes
003	82	Male	RE: late (neovascular) LE: -	RE: 8.5 LE: -	RE: 2/60 LE: 6/24+2	Diabetic retinopathy and cataract in both eyes
004	74	Female	RE: late (neovascular) LE: -	RE: 6 LE: -	RE: 6/7.5 LE: Hand movements	RE is the only eye Cataract in RE
005	83	Male	RE: no AMD LE: late (neovascular)	RE: - LE: 1	RE: 6/12 LE: 2/60	Glaucoma and cataract in both eyes
006	81	Female	RE: no AMD LE: late (neovascular)	RE: - LE: 0.5	RE: 6/48+2 LE: 6/7.5-2	Cataract in both eyes
007	76	Male	RE: no AMD LE: late (neovascular)	RE: - LE: 1	RE: 6/15-1 LE: 6/18+1	Glaucoma and cataract in both eyes
008	83	Female	RE: late (neovascular) LE: late (neovascular)	RE: 4.5 LE: 4	RE: 6/36-2 LE: 6/48+2	Cataract in both eyes
009	81	Female	RE: late (neovascular) LE: late (geographic atrophy)	RE: 10 LE: 1.5	RE: 6/60-4 LE: 6/15-1	Cataract in both eyes
010	75	Female	RE: late (neovascular) LE: no AMD	RE: 2	RE: 6/60-2 LE: 6/9-2	Cataract in both eyes
011	74	Female	RE: late (neovascular) LE: early	RE: 1 LE: 1	RE: 6/9+2 LE: 6/7.5-1	Cataract in both eyes; detached retina in RE
012	81	Female	RE: late (geographic	RE: 5 LE: 5	RE: 1/60 LE: 6/60-3	Cataract in both eyes

			atrophy and neovascular) LE: late (geographic atrophy)			
013	74	Male	RE: early LE: late (neovascular)	RE: 2 LE: 2	RE: 6/9.5+2 LE: 6/48-3	-
014	56	Male	RE: no AMD LE: late (neovascular)	RE: - LE: 1	RE: 6/7.5 LE: 6/24+2	Central serous retinopathy and cataract in both eyes
015	82	Female	RE: late (neovascular) LE: late (neovascular)	RE: 6.5 LE: 2	RE: 6/9+2 LE: 6/12-1	Cataract in both eyes
016	78	Female	RE: late (neovascular) LE: late (neovascular)	RE: 2 LE: 2	RE: 6/9-1 LE: 6/60-4	Cataract in both eyes
017	79	Male	RE: late (neovascular) LE: early	RE: 4.5 LE: 1.5	RE: 6/24+1 LE: 6/7.5	Cataract in both eyes
018	82	Male	RE: late (neovascular) LE: late (neovascular)	RE: 5.5 LE: 4.5	RE: 6/24 LE: 2/60	Glaucoma and cataract in both eyes; cystoid macular oedema in RE
019	66	Female	RE: early LE: late (neovascular)	RE: 2.5 LE: 2.5	RE: 6/18 LE: 6/15	Cataract in both eyes
020	77	Male	RE: late (neovascular) LE: late (neovascular)	RE: 9 LE: 7	RE: 6/12 LE: 6/7.5	Cataract in both eyes
021	87	Female	RE: late (neovascular) LE: late (neovascular)	RE: 2.5 LE: 2.5	RE: 6/60-2 LE: 6/48-1	Cataract in LE
022	82	Male	RE: no AMD LE: late (neovascular)	RE: - LE: 9.5	RE: 6/9-1 LE: 6/60-1	Polypoidal choroidal vasculopathy and cataract in both eyes
023	78	Male	RE: late (neovascular) LE: late (neovascular)	RE: 4 LE: 14.5	RE: 6/24 LE: 6/60-2	-
024	84	Female	RE: late (neovascular) LE: late (neovascular)	RE: 9 LE: newly diagnosed	RE: 6/7.5+2 LE: 6/9-1	Cataract in both eyes

025	59	Female	RE: late (neovascular) LE: -	RE: 3.5 LE: -	RE: 6/12 LE: 6/48	Myopic macular degeneration and cataract in both eyes
026	83	Female	RE: late (neovascular) LE: -	RE: 2 LE: -	RE: 6/36-1 LE: 6/6	Glaucoma in RE and cataract in both eyes
027	85	Female	RE: late (geographic atrophy and neovascular) LE: late (geographic atrophy)	RE: 6 LE: 6	RE: 6/30-2 LE: 6/60-3	Cataract in both eyes
028	77	Female	RE: early LE: early	RE: 6 LE: 6	RE: 6/12+1 LE: 6/48	Glaucoma and cataract in both eyes
029	72	Male	RE: late (geographic atrophy) LE: late (geographic atrophy)	RE: 7 LE: 7	RE: 6/15+2 LE: 6/60	Diabetic retinopathy in right eye

ID = Identification, AMD = Age-related macular degeneration, RE = right eye, LE = left eye

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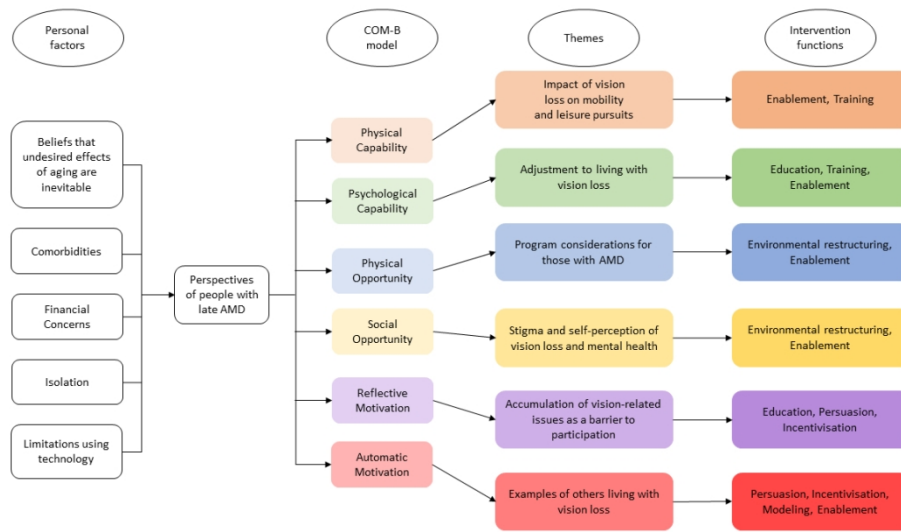


Figure 1. The impact of vision loss on mental health and mental wellbeing programs, from the perspectives of people primarily with late AMD, based on the Capability, Opportunity, Motivation and Behaviour (COM-B) model (inner wheel) and intervention functions (outer wheel) of the behaviour change wheel.²³ Personal factors which may have influenced these perspectives are also displayed.

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10 **Title:** Perspectives of people with late age-related macular degeneration on
11 mental health and mental wellbeing programmes: a qualitative study
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13 **Running head:** Mental health and age-related macular degeneration
14

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34 **Keywords:** mental health; macular degeneration; qualitative study; wellbeing
35 programmes; depression

36 **Author contributions:**
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10 **ABSTRACT**

11 **Purpose:** People with age-related macular degeneration (AMD) experience high rates of
12 depression, but rarely engage in or have access to tailored mental wellbeing programmes.
13 This qualitative study investigated the perspectives of those primarily with late AMD on
14 mental health and mental wellbeing programmes.
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17 **Methods:** Twenty-eight people with late AMD in at least one eye, and one person with early
18 AMD in both eyes, aged 56-87 years (mean age 78 years) attending a private eye clinic
19 between December 2019 and January 2020 in Sydney, New South Wales, Australia,
20 participated. Individual semi-structured interviews were conducted and analysed deductively
21 using content analysis, following the individual level factors for health promotion
22 interventions in the behaviour change wheel: *Capability (Physical & Psychological)*,
23 *Opportunity (Physical & Social)*, and *Motivation (Reflective & Automatic)*.
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26 **Results:** Six major themes were identified: *Capability*: (1) Impact of vision loss on mobility
27 and leisure pursuits; (2) Adjustment to living with vision loss; *Opportunity*: (3) Program
28 considerations for those with AMD; (4) Stigma and self-perception of vision loss and mental
29 health; *Motivation*: (5) Accumulation of vision-related issues as a barrier to participation; (6)
30 Examples of others living with vision loss. General personal factors relevant to delivery of a
31 program in this age group were also identified: Comorbidities; Limitations using technology;
32 Isolation; Financial concerns; and Beliefs that undesired effects of aging are inevitable.
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36 **Conclusions:** Complex individual, environmental and social factors influence the
37 perspectives of people with late AMD on mental health, and potential participation in mental
38 wellbeing programmes. These factors should be considered when developing and
39 implementing mental wellbeing programmes to improve the emotional and functional
40 rehabilitation outcomes for people with AMD.
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INTRODUCTION

Age-related macular degeneration (AMD) is a progressive, chronic disease impacting central vision, and is a major cause of low vision and blindness in older age.^{1,2} Globally, 196 million people were affected by AMD in 2020, with this number projected to increase to 288 million by 2040.³ The two main stages of AMD are early and late, with vision loss characteristic of late AMD. Late AMD is further subdivided into neovascular and geographic atrophy AMD, with the classification based upon the presence or absence of choroidal neovascularisation (abnormal blood vessels), respectively.⁴ The neovascular form affects around 10-15% of people with AMD and accounts for approximately 90% of cases of severe vision loss due to AMD.⁵ Neovascular AMD can be managed with a range of antiangiogenic treatments but there is often persistent vision impairment.⁶

Despite treatment, AMD can have a profoundly negative effect on the quality of life. The greatest effects have been reported as a result of restrictions in reading, leisure activities and shopping.⁷ Irrespective of the degree of vision loss, there is also a high level of emotional distress, such as worry and frustration about vision loss, with many expressing concern about their eyesight getting worse and their ability to cope with everyday life.⁷ Notably, ratings of quality of life and emotional distress have been shown to be significantly worse among people with AMD, compared with similarly aged community dwelling adults with chronic illness.⁸

The prevalence of depression among patients with AMD is 32.5% (n=151, >60 years old),⁹ i.e., twice the frequency identified in studies of community-dwelling older people.^{10,11} This is very concerning given that 33% (n=51) of those with depression also have greater levels of vision-specific and general physical disability, and slightly worse visual acuity than non-depressed participants.¹² This highlights the close relationship between AMD and mental health disorders, with both cumulatively leading to substantial functional impairment.

Although mental health disorders are treatable, approximately 60% of a nationally representative sample of Australians aged 55 to 64 years meeting the criteria for a mental disorder did not make use of mental health services.¹³ Similar data are seen in American older adults, where approximately 70% (n=348, >55 years old) with mood and anxiety disorders did not use mental health services.¹⁴ This highlights the need for and access to programmes that focus on mental wellbeing for older people, particularly those with AMD who are at risk of the cumulative impact of vision loss and mental health issues. Shared-

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decision making has been suggested to improve adherence to mental health services and patient-centred care.¹⁵ This serves to heighten trust and acceptance of potential evidence-based programmes ensuring adherence and mental health issues experienced by this population can be appropriately targeted with effective interventions.

Low-vision rehabilitation services have been shown to improve vision-specific quality of life; however, these services were shown to have little impact on depressive symptoms.¹⁶ A low-vision self-management programme has also been shown to have very little benefit on quality of life in older adults accessing low vision services.¹⁷ However, psychosocial interventions¹⁸ and psychological therapies¹⁹ have been shown to have a positive impact on depression in adults aged 18 years or older with low vision. These psychological therapies have been mainly developed and tested in people with AMD, and to a lesser extent in those with glaucoma, Stargardt disease, or in general low vision or unspecified populations.¹⁹ We aimed to investigate the perspectives of people with AMD on mental wellbeing programmes in a novel qualitative study in order to involve them meaningfully in planning, developing and implementing a psychological programme targeting depression and anxiety. Given the high prevalence of AMD and the debilitating nature of the condition itself and associated issues such as elevated anxiety and depression, the development of interventions targeted to this population is integral to improving their quality of life and functional independence.

METHODS

This study is reported in line with the Consolidated Criteria for Reporting Qualitative Research (COREQ) statement (Supplementary File 1).²⁰

Design and setting

A qualitative study investigated the perspectives of people primarily with late AMD and the factors that influenced their participation in mental wellbeing programmes. Semi-structured interviews were used due to their flexibility and ability to collect rich narrative data.²¹ This study was conducted at the Sydney West Retina Clinic (private eye clinic), located in Westmead, Sydney, Australia. Patients of this high-volume clinic are not required to pay any out of pocket costs if eligible for Medicare rebates.

Study participants

People with AMD were recruited from the clinic using convenience sampling, and recruitment stopped when data was considered saturated. Data saturation was assessed at regular weekly project meetings by researchers (LD, SG), and considered achieved when

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additional data did not lead to the development of new codes.²² People diagnosed with AMD were approached by researchers (DT, SG) during their visit to the clinic, and invited to participate. The inclusion criteria were: (1) diagnosed with AMD (both early and late) and (2) aged 50 years or older. Interviewers explained the purpose of the study and gained verbal informed consent before participation, ensuring involvement was voluntary.

Data collection

One-to-one, face-to-face, semi-structured interviews were conducted with consenting participants who met the inclusion criteria. Interviews were conducted by an undergraduate medical student (SG), who received initial and ongoing training from an experienced Faculty of Medicine and Health PhD student (DT), and two senior qualitative researchers (BG, LD). A semi-structured guide was developed using the behaviour change wheel framework.²³ The framework emphasises the relationships between individual behaviour change and intervention design. This was appropriate to understanding the perspectives of people with AMD about mental health and consequent design of a mental health programme.

The interviews were performed in a quiet room at the clinic, but could be interrupted when a participant was called on by a clinician for treatment or consultation, and then resumed when possible. The questions were focused on exploring the experience of participants and what support or programmes they would like to engage with, should these be available. The open-ended question style served to facilitate more responses from participants, and allowed them to expand on and discuss their thoughts.²¹ The semi-structured interview guide is included in Supplementary File 2. Some examples of specific questions included; “Can you tell me a bit more about your macular degeneration?”, “What is your understanding regarding support programmes for AMD patients to help cope with the condition?”, “Are you aware of any support services out there in general?”, “What are some concerns you have regarding your mental health?” and “What do you think should be involved in a mental health programme developed for people with vision impairment?”.

Each interview was audio recorded, transcribed verbatim and analysed using word processing software. Transcripts were not returned to participants for comment and/or correction, nor were repeat interviews conducted. Participant demographic information was collected from the medical record at the clinic following completion of each interview. This included age (years), sex (male/female), AMD type (late/early) in each eye, time since AMD diagnoses

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10 (years), visual acuity in each eye, and history of other eye conditions. Some participants were
11 initially seen or treated at another eye clinic, and consequently not all data were available.
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13 **Data analysis**

14 The qualitative data were analysed using the framework from the behaviour change wheel.²³
15 Specifically, data were deductively coded using content analysis,²⁴ under the headings of the
16 sources of behaviour (Capability, Opportunity, Motivation and Behaviour (COM-B) model,
17 inner wheel), and the intervention functions (middle wheel).²³ Due to broad application of the
18 behaviour change wheel as a framework, it was not expected that all sources of behaviour or
19 intervention functions would be found to be relevant. Only data that were specifically related
20 to the impact of AMD on the psychological functioning of participants, and perspectives
21 related to the design of a potential programme, were analysed. Personal factors, which may
22 have not been a direct result of vision loss but would likely influence perceptions and
23 participation in a programme, were also documented. We took an iterative approach to data
24 analysis,²⁵ whereby data were revisited, coded, and themes discussed many times to ensure
25 analytic reflexivity. Transcripts were coded separately by two researchers (SG, LD), who met
26 weekly to reflect on coding and theme development after each set of 10 transcripts were
27 completely read and coded. Initial themes were agreed upon, along with their illustrative
28 quotes, and presented at two meetings with a wider group (BG, LK, DT, SG, LD) for critique
29 and finalisation.
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36 **Patient and public involvement**

37 No participants or members of the public were involved in the design, recruitment or conduct
38 of the study. All participants were informed via email or telephone call about the publication
39 of the results.
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41 **Ethics approval**

42 Ethics approval was granted by the University of New South Wales Human Research Ethics
43 Committee (HC190356).
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46 **RESULTS**

47 **Participant characteristics**

48 Twenty-nine participants with AMD were recruited from December 2019 to January 2020. A
49 further participant had been recruited, but their vision impairment did not include a diagnosis
50 of AMD, and they were subsequently excluded. Average age was 78 (standard deviation = 7,
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10 range = 56 to 87) years old. Participants were mostly female (16/29, 55%), and had late stage
11 AMD (28/29, 97%) in at least one eye (Table 1).

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13 Interviews lasted an average of 17 minutes (range = 4 to 50 minutes) and varied in length
14 depending on the depth of answers.

15 16 **Themes**

17 Six major themes were identified which corresponded with all elements of the Capability,
18 Opportunity, Motivation and Behaviour (COM-B) model, and seven of the nine intervention
19 functions of the behaviour change wheel.²³ The themes and their relationship to the behaviour
20 change wheel are summarised in Figure 1, and quotes that are more illustrative are presented
21 in Supplementary File 3.

22 23 ***Physical Capability (Training, Enablement)***

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25 *Theme 1: Impact of vision loss on mobility and leisure pursuits.* Participants reported that
26 vision loss had greatly affected their mobility, and this lack of mobility did not just reduce
27 access to support, but also led to increased isolation from the community, friends and family:

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31 ‘What am I doing now? Sitting on my bum all bloody day because I can’t do
32 anything else. I can’t see to go out, I can’t catch a bus. I can’t catch a train, you
33 know?’

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35 Vision loss reduced their ability to drive, or ability to drive at certain times or for long
36 distances. For some, this changed their perception of themselves as independent, competent
37 individuals:

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41 ‘I used to do a lot of driving. I’ve been driving for approximately 65 years. I used
42 to work for a big company. I used to travel interstate all the time...I don’t drive
43 anymore. I don’t trust myself anymore.’

44 In particular, all participants mentioned how their vision loss reduced their ability to read:

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47 ‘I used to be a very, very avid reader of books. I am no longer an avid reader of
48 books.’

49 Most participants also reported that the vision loss associated with AMD led to a loss of
50 ability to engage in hobby and leisure activities, which was sometimes conceptualised as an
51 inability to participate in all activities:

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10 'I used to do a lot of crosswords and I used to play a lot of tennis – I can't do
11 anything now.'

12
13 However, those who could get access to a low vision aid appeared to be able to continue
14 reading and engage in hobby and leisure activities:

15
16 'I'm still able to read but I have a special magnifying glass. It's a good one too.
17 Smaller print, big print – I'm okay.'

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19
20 ***Psychological Capability (Education, Training, Enablement)***

21 *Theme 2: Adjustment to living with vision loss.* Participants reported varying levels of
22 adjustment to their vision loss. This was mediated by their outlook on diagnosis, acceptance
23 of vision loss and fear of further vision loss:

24
25 'I was a pretty secure sort of person. I suppose I don't feel as secure as I used to. I
26 take a risk and go, I won't now because I fear I will fall, or I'll walk into – it's
27 amazing when you can't see properly how much it affects your whole life.'

28
29 'And for me, at the moment, I don't consider it a disability. Because I can still see
30 with one good eye. Because I can drive, I can still read, I can use my laptop.'

31
32
33 The impact of vision loss on confidence stopped some participants from getting out in their
34 community with others, despite having the opportunity to do so:

35
36 'Your confidence, for one thing. I do get asked out a lot. I've got nice friends
37 from the church and all that but I think no, I won't go cause they've got to watch
38 me and it's only – I'm the one that's stopping myself, not my friends, they are all
39 eager to come and take me out to whatever's on. It's hard when you've been
40 independent and then this happens.'

41
42 'It's just amazing that it [vision loss] makes you very fearful.'

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46 ***Physical Opportunity (Environmental restructuring, Enablement)***

47 *Theme 3: Programme considerations for those with AMD.* None of the participants
48 interviewed were aware of mental wellbeing programmes designed for people with vision
49 impairment or AMD:

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51 'I haven't heard of anything as such around our area. I mean there might be stuff,
52 I just haven't...heard of it, no.'

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10 Participants offered that more appropriate and increased advertising would improve
11 awareness of existing and future support programmes:
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13 'I just think we should advertise the help that we can get more because I don't
14 know what help we've got, but like my son keeps saying, there must be help for
15 people like you. And I'm willing to try things where some might not. I've given
16 up. I mean, I've been like this two year[s]. I've sort of thought I'll just get on and
17 make my own life, yes, so that's what I've done.'
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20 Participants also reported concerns regarding the location, cost and accessibility of existing
21 and proposed programmes:
22

23 'But there's just really nothing within an area, easiness to get to [is important].
24 So, it can be a bit difficult.'
25

26 'You also need to think those with visual problems probably don't have the
27 money to come together easily in groups.'
28
29

30 'It [community programme] was no good to me. They would write things on the
31 blackboard and you've got to tell them what it is. Well I couldn't see those
32 things.'
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35 Participants however recognised the benefits of attending programmes on their wellbeing:
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37 'I think it's good for your morale. And I think if you're even people that [sic] are
38 semi-blind or totally blind, they come along and join in with these functions. It's
39 a good feel...one of the old ladies saying that she usually sleeps every afternoon.
40 That particular day, she decided to come to the function, and she enjoyed it so
41 much.'
42
43

44 ***Social Opportunity (Environmental restructuring, Enablement)***

45 *Theme 4: Stigma and self-perception of vision loss and mental health.* Some considered their
46 vision loss or mental health problems were not severe enough to seek services, especially
47 when compared with other people:
48

49 'People with vision impairment are very depressed. I mean, to be honest, it's
50 depressing. But as far as I'm concerned, being depressed is something that you
51 have to work your way through or compensate for or whatever.'
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10 'My GP wants a form filled in for depression and I laughed at her. She's more
11 depressed than me.'
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13 However, many of the participants saw social connection with others, including those with
14 vision impairment, as positive:
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16 'I like it, but a lot of the activities are more of a social event. And I look forward
17 – I like having – I wake up in the morning and if there's an event on today that I
18 really want to go to...and enjoy it...we have our happy hours and function days.'
19

20
21 'And then afterwards we sit down, and we have a chat and that is more important
22 for me that I can talk to someone. More important than anything else.'
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25 ***Reflective Motivation (Education, Persuasion, Incentivisation)***

26 *Theme 5: Accumulation of vision-related issues as a barrier to participation.* Participants
27 were less motivated to engage in a programme due to an accumulation of practical and
28 psychosocial concerns related to their vision loss.
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31 'If you talk to people who are vision impaired, it means they also have to have
32 somebody who's able to take them, bring them home. And it becomes a big
33 effort...[the support] person themselves may not want to give up half a day.'
34

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36 'Just once you get it, you get it and that's it, sort of thing. See it's all like the girl
37 or the woman, she used to take me out for a walk every so often. Well what's the
38 good of that, I can't see things properly and sometimes I trip over things and, you
39 know, it's just a waste of time sort of thing...and they say I can go out in their
40 bus on outings, but what's the good of that? I can't see properly.
41

42 This led to loss of hope or lack of motivation to try anything new or access existing services:
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44 'I had a girl from the council, I think it was from the council, but she tried to help
45 me but there's nothing you can do really.'
46

47
48 ***Automatic Motivation (Persuasion, Incentivisation, Modelling, Enablement)***

49 *Theme 6: Examples of others living with vision loss.* Participants would commonly refer to
50 how others were living or coping with their vision loss, negatively and positively:
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10 'One of my friends she sort of...let[s] people know that she has low vision and
11 that. And she just went on a cruise with her family recently. She said it was very
12 good because...people were very sort of helpful to her.'

13
14 'One woman there...she's about 95...she said that [she] didn't help with the
15 macular at all. She's had them for years. She's still quite good in herself. She's
16 restricted so much with the lack of vision. Well, she always seems to be in good
17 spirits, I always say g'day to her when I see her in the morning. If I see her
18 coming, I actually hold the chair out for her and help her into a chair.'

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21
22 Participants highlighted the need for good leadership for the support groups, and the
23 importance of having a positive experience the first time they attend:
24

25 'That's because of the staff. That's what made a difference. That was the key.'

26
27 'You always need leadership. Yeah, and nice leadership, open-style
28 leadership...if you can find someone important enough or it's – preferably a lady
29 that would be followed by women, because women tend to lead their social
30 circle.'

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33 'You only need to meet one person that you can connect with and you want to go
34 back there.'

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36 'You've got to make sure the first time they attend, it's got to be a good show,
37 otherwise they'll think, that was a waste of time, I'm not coming back... first
38 impressions are always very important.'

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41 **Personal factors**

42 Interviews also revealed complex individual, environmental and social factors influencing
43 participants' perspectives. Although these personal factors, or their impact, were likely
44 exacerbated by vision loss, they are also commonly experienced by older adults more
45 generally. These factors were readily reported by many of the participants and are likely to
46 affect how people primarily with late AMD engage in mental wellbeing programmes. The
47 personal factors are presented in Figure 1, and illustrative quotes are presented in
48 Supplementary File 4.
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10 **DISCUSSION**

11 This study explored the perspectives of people primarily with late AMD regarding mental
12 health and wellbeing programmes, and considered the individual and interventional factors
13 that influence engagement in such programmes. Of note, most participants were unaware of
14 existing general low vision rehabilitation programmes. Further, none of the participants were
15 aware of programmes specifically aimed at improving the mental wellbeing of those with
16 AMD. Past research has demonstrated lack of knowledge as a major barrier to receipt of
17 mental health services in older adults with vision impairment (n=871, mean age=73 years).²⁶
18 This indicates a worryingly large gap between available services and knowledge given the
19 high frequency of mental health concerns, isolation and loss of functional independence
20 among people with AMD.^{9, 12} Further, older adults with vision impairment who also have
21 depression are more vulnerable to further health decline.²⁷ However, participants provided
22 critical insights into the barriers preventing them from engaging in mental wellbeing
23 programmes, if they were available, and offered suggestions as to what elements they would
24 look for in a programme and how these could be implemented. In particular, participants
25 expressed a desire for increased social connection to ameliorate feelings of isolation or
26 loneliness.
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33 *Adjustment to living with vision loss* was a major theme identified under *psychological*
34 *capability*. Our participants had varied outlooks on their vision loss and AMD diagnosis
35 including denial, fear of further decline and acceptance of living with vision loss. We
36 observed better functioning in those with social support, longer-term vision loss, those in
37 receipt of low vision rehabilitation services and those with access to low vision aids.
38 Conversely, we observed worse functioning in those with a fatalistic attitude to the
39 undesirable effects of aging, those fearful of further vision loss and those isolated or not in
40 receipt of low vision rehabilitation services. Adjustment and/or acceptance of vision loss has
41 been shown to increase psychosocial wellbeing and use of services.²⁸ However, less than
42 20% of older adults with vision impairment engage in rehabilitation services.²⁹ Further, those
43 with recent vision loss are more likely to experience higher levels of psychological distress,
44 compared to those with longer-term loss.³⁰ Although a lack of adjustment to vision loss may
45 explain why many of our participants were not in receipt of services, our findings suggest
46 that many were completely unaware of existing services. *Education* may be a mechanism to
47 influence *psychological capability*, as well as *reflective motivation* to engage in a
48 programme, indicating people should be provided with education early in their diagnosis of
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AMD. With appropriate training and resources, vision rehabilitation^{31, 32} or eye care professionals³³ may be well placed to deliver this education.

Stigma was a major theme identified under social opportunity, and impacts how those with vision impairment cope with their mental health and vision loss,³⁴ and their uptake of psychological support.³⁵ Stigma is a major inhibitor of social well-being in older adults with vision impairment,³⁶ who often find it difficult to speak with their physician about their concerns.³⁷ Vision impairment is considered a challenge to identity, with recognition of vision impairment akin to being 'devalued' or 'sub-human'. Consequently, those with vision impairment were less likely to use mobility aids or engage in social activities in an attempt not to draw attention to their vision impairment.³⁶ Similarly a diagnosis of AMD can lead to isolation if individuals chose not to disclose their condition to others.³⁸ Programmes aimed at reducing stigma by taking a holistic approach to supporting those adjusting to living with AMD have been recommended. *Education, Enablement and Modelling*²³ and associated behaviour change techniques³⁹ could be used to design a programme to reduce stigma, and enhance the psychosocial wellbeing of people with AMD.

Our themes of *Impact of vision loss on mobility and leisure pursuits* and *Accumulation of vision-related related issues as a barrier to participation* were similar to themes identified in other qualitative studies in people with AMD: functional limitations, adaptation and independence.³⁸ Vision loss due to AMD has previously been found to result in limitations of activities of daily living, including reading and driving,³⁸ particularly for those with bilateral vision loss. There is vast scope for improved access to and facilitated engagement in low vision rehabilitation services and mental wellbeing programmes, which may lead to a continuation of, or improvement in usual daily activities.

Strengths and limitations

Semi-structured interviews allowed for tailored discussions and a deeper understanding of the impact of AMD on mental health and perspectives on mental wellbeing programmes. Data saturation was achieved, and we are confident the themes captured the perspectives of this group. However, the study is limited to people who attended one ophthalmology clinic in New South Wales, Australia, the majority of whom have late AMD. Patients of the clinic are not required to pay any out of pocket costs, and see the same healthcare practitioners, reducing the representativeness of our population. Their opinions may be different from people with early AMD, or those who are not receiving treatment or presenting to a different

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clinic. Recruitment and completion of interviews in a quiet room at the clinic was convenient for the participants. However, the flow of some of the interviews were likely interrupted when a participant was called away by the clinician for treatment or consultation during the interview, which may have led to shorter interviews. Additionally, access to definitive information regarding duration and severity of vision loss may have improved our understanding of the impact of AMD and adjustment over time, as well as establishing potential differences in perspectives of those with milder versus more severe vision loss. Finally, none of the participants were aware of mental wellbeing programmes, which may have been a result of the convenience sampling used. Future studies should look to use purposeful sampling of patients with AMD who have previously engaged in a mental wellbeing programme.

Implications for practice and research

We found that adjustment to vision loss due to AMD, particularly in those with late AMD, is different for every person. Our findings suggest that in order to facilitate adjustment, programmes should aim to reduce the stigma associated with vision loss, account for individual circumstances, and promote engagement in current services. Accessibility is also critical to helping people with AMD participate in appropriate programmes, including regular long-term involvement. Accessibility not only includes accounting for the unique travel concerns faced by those with AMD, but also appropriate marketing, ensuring programmes are engaging by using meaningful speakers and activities and offer opportunities for participants to have some ownership over their content and delivery. Any programmes developed will need to be rigorously tested to ensure efficacy before widespread implementation.

Acknowledgements: This study was supported by the UNSW SOVS Macular Degeneration Research Grant from the Estate of the late Peter Anthony John Vild. We thank the participants that took part in this research.

Disclosure: The authors declare no conflicts of interest and have no proprietary interest in any of the materials mentioned in this article.

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Figure 1. The impact of vision loss on mental health and mental wellbeing programmes, from the perspectives of people primarily with late age-related macular degeneration (AMD), based on the Capability, Opportunity, Motivation and Behaviour (COM-B) model (inner wheel) and intervention functions (outer wheel) of the behaviour change wheel.²³ Personal factors, which may have influenced these perspectives, are also displayed.

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