

# SESSION 2: Why the Eye: Forum on the Public Health Connection Between Mental Wellness and Vision





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# INTRODUCTION TO: WHY THE EYE: FORUM ON THE PUBLIC HEALTH CONNECTION BETWEEN MENTAL WELLNESS AND VISION

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### EFFECTS OF VISION LOSS

- People with vision loss are more likely to experience:
  - Depression
  - Diabetes
  - Hearing impairment
  - Stroke
  - Falls
  - Cognitive decline
  - Premature death
- Decreased ability to see is associated with reduced ability to:
  - Drive
  - Read
  - Manage finances
  - Travel
- The cost of vision loss, including direct costs and lost productivity, is estimated to exceed \$35 billion (Rein, Zhang, Wirth, et al., 2006)

- When people first access vision rehabilitation services, data from the USA indicate that the prevalence of depression is ~ 30%.
  - Brody, et al., 2001, Depression, visual acuity, comorbidity, and disability associated with age-related macular degeneration. Ophthalmology;
  - Brody, et al., 2005, Self-management of age-related macular degeneration at the 6-month follow-up - A randomized controlled trial. Arch Ophthalmol.
  - Rovner, et al., 2001, Neuroticism predicts depression and disability in agerelated macular degeneration. J Am Geriatr Soc.
  - Rovner, et al., 1997. Screening for depression in low-vision elderly. Int J Geriatr Psychiatry.
  - Rovner, et al., 2007. Preventing depression in age-related macular degeneration. Arch Gen Psychiatry.

### INTERNATIONAL FINDINGS

COUNTRY	RATE OF DEPRESSION	
Thailand	43% (during COVID)	
Bangladesh	58% (during COVID)	
Brazil	54% (during COVID)	
Australia (pts. w/ cataracts)	31%	
England (older adults with VI)	30%	
Netherlands (older adults with VI)	27%	
India (older adults with VI)	21%	

Summary from Tantirattanakulchai et al., 2023, Clinical Ophthalmology

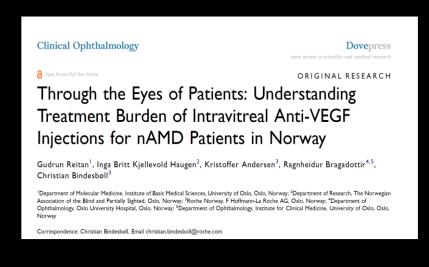
# OVER 1/3 OF PEOPLE BELIEVE THEY NEED EMOTIONAL HELP AND COUNSELING AT THE TIME THEY BEGIN RECEIVING SERVICES FOR B/VI

- "If I had someone to talk with about my loss of sight; I would have appreciated psychological help to discuss emotional problems, such as anger and grief. Right now I would still like and appreciate this help" (Conyers, 1992)
- ~25% of people said they felt unable to talk to people in their life about their concerns (Conyers, 1992)

### CHANGES IN EMOTIONAL REACTIONS FROM TIME OF LOSS OF SIGHT TO UP TO 2 YEARS LATER (N=104) (FROM CONYERS, 1992)

REACTION	<u>IMPROVED</u>	DECLINED
Nothing to get up for	0	17
Nothing to look forward to Meaning and purpose left life	3 2	26 23
No hope for the future Bitter about loss of sight	3 2	26 24
Felt sad	3	34
Felt I would not feel again Couldn't accept it happened to me	1 3	11 35
Could not accept sight would not	4	26
return Felt embarrassed	0 2	28 28
Felt so alone in this experience	2	11
Felt like a second class person	2	15

# ASPECTS OF **TREATMENT** CAN FURTHER IMPACT MENTAL HEALTH



- 130 nAMD patients responded to the survey. The majority of patients reported to receive nine or more injections per year.
- Many patients needed caregiver support for every treatment appointment (37.7%).
- The week before the treatment, 33.1% of patients reported to be stressed and 15.4% struggled to sleep.
- The majority of patients reported the treatment as uncomfortable (54.6%) or as somewhat painful (26.2%).

# PEOPLE WITH BLINDNESS OR LOW VISION FREQUENTLY ENCOUNTER STIGMATIZATION IN THEIR DAILY LIVES



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#### Coping With Stigmatization and Discrimination Related to Blindness and Low Vision

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Purpose/Objective: Research suggests that effective coping with stress related to stigmatization and discrimination might protect the stigmatized persons against poor mental health outcomes associated with experiencing stress. Despite this, research specifically on coping with social stigma among people who are blind or have low vision has received less attention. We addressed this gap in the literature, by exploring the lived experiences of people who are blind or have low vision with regard to their perceptions of stigma and their coping responses. Research Method/Design: We employed a qualitative research method using an Interpretative Phenomenological Analysis (IPA). Participants were 11 Australians who were blind or had low vision, aged between 18 to 65 years, and were recruited via advertisement. Data was collected using in-depth interviews and was subjected to data analysis using IPA. Results: Two sets of themes emerged from the data analysis which we classified under 2 superordinate themes labeled as Ways of Coping and Coping Skills and Resources. Conclusions/Implications: This study highlights the significance of personal and contextual aspects of coping with stigma and discrimination by demonstrating the complex nature of how people who are blind or have low vision cope with stigma. Coping efforts might be tenuous and require cognitive and/or behavioral skills and psychological resources and as such, individuals experiencing stigma might benefit from interventions aimed at improving coping skills and resources, such as self-advocacy, assertion, or positive reinterpretation.

- This can lead to:
  - not participating in rehabilitation programs (Southall & Wittich, 2012)
  - limited social inclusion (Bennion et al., 2012; Kong & Loi, 2017)
  - a reduced sense of mastery
  - anger
  - feeling uncomfortable in varied social interactions
  - shame and embarrassment (J. Green et al., 2002; Nyman et al., 2012; Thombs & de Board, 2016).

### STAGES OF PSYCHOLOGICAL ADJUSTMENT TO LOSS OF SIGHT

- From Cholden, 1958, A Psychiatrist Works With Blindness
  - Depersonalization
  - Depression
  - Recovery
- From Tuttle & Tuttle, 2004, Self-Esteem and Adjusting with Blindness\*
  - Trauma, physical or social
  - Shock and denial
  - Mourning and withdrawal
  - Succumbing and depression
  - Reassesment and reaffirmation
  - Coping and mobilization
  - Self-acceptance and self-esteem

Reaction

Rebuilding



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REVIEW

#### Visual Impairment and Mental Health: Unmet Needs and Treatment Options

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associated with a substantial economic burden, reduced quality-of-life, concurrent medical issues, and mental health problems. In this review, the mental health needs of people with vision impairment are examined. Patients and methods: A review of recent literature on mental health outcomes and

**Purpose:** An estimated 2.2 billion people are visually impaired worldwide. Given that age-

related vision loss is a primary cause of vision impairment, this number is projected to rise with increases in average lifespan. Vision loss often results in significant disability and is

current treatments in people with visual impairment was conducted.

**Results:** Considerable data indicate that rates of depression and anxiety are elevated among people with visual impairments. Moreover, individuals of lower socioeconomic status may be at increased risk for vision impairment and subsequent mental health problems. Existing psychosocial interventions for improving mental health in people with visual impairment show some promise, but are limited by low adherence and lack generalizability.

Conclusion: In order to improve outcomes, a better understanding of the mechanisms linking visual impairment and poor mental health is needed. It will also be essential to develop more effective interventions and expand access to services to improve the detection and treatment of mental health problems in this population.

**Keywords:** visual impairment, vision loss, mental health, psychosocial interventions

https://www.ncbi.nlm.nih.go v/pmc/articles/PMC7721280/

### **SESSION 2: Why the Eye: Forum on the Public Health Connection Between Mental Wellness and Vision**



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### Patient Perspective



### **Carol Johnson**

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