The Michigan Vision-related Anxiety Questionnaire

INNOVATION PARTNERSHIPS

TECHNOLOGY NUMBER: 2021-206





OVERVIEW

- Validated Patient Reported Outcomes (PRO) measure
- Focused on psychosocial screening and monitoring for patients with inherited retinaldegenerations
- Measure can be utilized in therapeutic clinical trials for measuring the benefit of an investigational therapy on a patient's vision related anxiety

INNOVATION

An important new feature of the MVAQ is its ability to relate psychosocial reports to a retinal physiologic pathway. Furthermore, the MVAQ disability scores exhibit correlations with VA, phenotype, and self-reported feelings. Other PRO measures ask about a patient's feelings or concerns, but their responses cannot be linked to the pathologic defect underlying their visual impairment. In contrast, the MVAQ allows a clinician to map the concerns of the patient into an actionable area for improvement. As opposed to knowing that a patient experiences "fear" or "worry", with the MVAQ a clinician can identify whether a patient's rod dysfunction is the predominant cause of their anxiety, and recommend accommodations to their living setting and vision-related activities to address these specific anxiety-provoking situations.

The current study demonstrates psychometric validation of items pertaining to vision related anxiety in patients with IRDs and collectively form the MVAQ. The MVAQ measures visionrelated anxiety due to dysfunction in either a predominantly cone pathway or a rod pathway. It can be used as a screening tool to direct the attention of a clinician to counseling and psychosocial interventions, as well as to measure baseline and follow-up vision related anxiety for targeted low vision rehabilitation and psychotherapy. Finally, the MVAQ may also be **Technology ID** 2021-206

Category Content

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incorporated in therapeutic clinical trials to understand the benefit of a novel therapeutic intervention for reducing a patient's vision-related anxiety.

OTHER