

# Impact of Herpes Zoster Ophthalmicus on Healthcare Utilization and Patient Reported Outcomes: Results from a Multicenter Cohort Study

Laura T. Pizzi<sup>1</sup>, Benjamin E. Leiby<sup>2</sup>, Ayako Shimada<sup>2</sup>, Herbert J. Ingraham<sup>3</sup>, Emily W. Gower<sup>4</sup>, Joseph P. Shovlin<sup>5</sup>, Beth R. Friedland<sup>6</sup>, Haresh Ailani<sup>7</sup>, Katherine M. Prioli<sup>1</sup>, **Kejsi Begaj<sup>1</sup>**, Alin Kalayjian<sup>1</sup>, Justin D. Gatwood<sup>8</sup>, Ann P. Murchison<sup>9</sup>

<sup>1</sup>Rutgers University, Piscataway, NJ, USA; <sup>2</sup>Thomas Jefferson University, Philadelphia, PA, USA; <sup>3</sup>Geisinger Health System, Danville, PA, USA; <sup>4</sup>Gillings School of Global Public Health, University of North Carolina at Chapel Hill, NC, USA; <sup>5</sup>Northeastern Eye Institute, Scranton, PA, USA; <sup>6</sup>UNC School of Medicine, Chapel Hill, NC, USA; <sup>7</sup>Eye Consultants of Northern Virginia, Springfield, VA, USA; <sup>8</sup>GSK, Philadelphia, PA, USA; <sup>9</sup>Wills Eye Hospital, Philadelphia, PA, USA

## Background

- Herpes zoster, commonly known as shingles, **can lead to herpes zoster ophthalmicus (HZO)**, which affects the eye and surrounding areas. **The frequency of HZO out of all herpes zoster cases in the United States (US) is 4.2% and ranges from 2.7% to 6.7% annually**, with older adults and immunocompromised individuals at higher risk.<sup>1,2</sup>
- HZO is a severe manifestation of herpes zoster happening when the ophthalmic branch of the trigeminal nerve is involved, and it could lead to **severe complications**. These complications often necessitate **long-term medical treatments** and frequent **follow-up evaluations**.<sup>3</sup>
- Patients with HZO often experience a **diminished quality of life** due to persistent symptoms and the need for lifestyle modifications. The chronic nature of the disease can also lead to significant **depressive symptoms**.<sup>4,5</sup>



The primary objective of this study was to **assess the burden of HZO in the US** by evaluating the **frequency and duration of specific complications**, the **prevalence of depressive symptoms**, the **impact on health-related quality of life**, and **healthcare resource utilization**.

## Methods

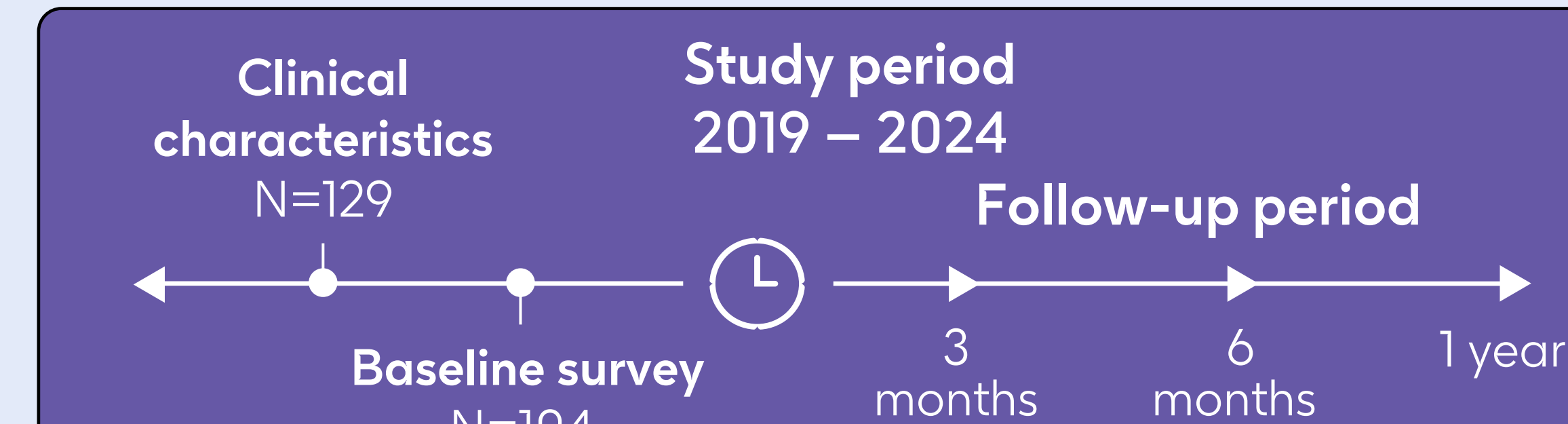
### Real world observational prospective cohort study

#### ✓ Patient eligibility criteria

- Active clinical diagnosis of HZO (initial or recurrent)
- ≥18 years of age
- English or Spanish speaking

#### ✗ Patient exclusion criteria

- Simultaneous enrollment in a clinical trial focused on HZO



#### Dataset

- 6 US ophthalmologic practices**  
Eye Consultants of Northern Virginia, Geisinger Health System, Metropolitan Eye Research and Surgery Institute, Northeastern Eye Institute, University of North Carolina, Wills Eye Hospital
- Total patients:** 130 consented (one patient did not complete the clinical information form)
- Complete data:** 110 completed at least 1 survey

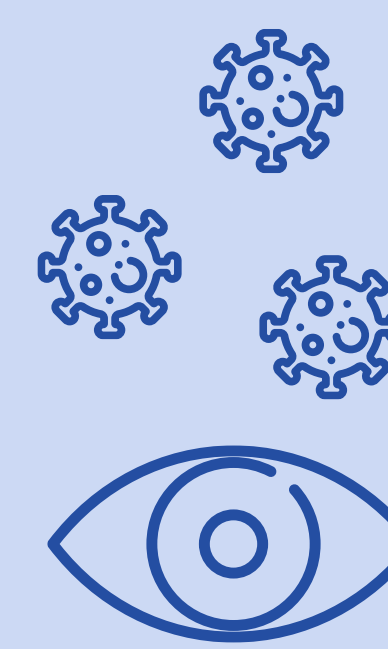
#### Validated patient-reported study instruments

- Eight-item Patient Health Questionnaire (**PHQ-8**) for assessment of severity of depressive symptoms<sup>6</sup>
- National Eye Institute 25-Item Visual Function Questionnaire (**NEI-VFQ-25**) for assessment of visual functioning (can be scored to health utilities)<sup>7,8</sup>
- Zoster Brief Pain Inventory (ZBPI) for assessment of pain items<sup>9</sup>
- Work Productivity and Activity Impairment-Specific Health Problem questionnaire (**WPAI-SHP**) for assessment of the impact of health problems on work productivity (presenteeism and absenteeism)<sup>10</sup>

#### Analysis

- Descriptive statistics
- Generalized linear mixed models examined changes in outcomes over time for healthcare utilization and PHQ-8

## Conclusions



HZO patients experienced **significant clinical symptoms** and required **substantial healthcare resources**, especially within the first three months post-onset.



HZO significantly affected patients' **quality of life and mental health**, with long-term impacts on vision-related quality of life despite a decrease in depressive symptoms and healthcare utilization over time.



Findings highlight the **need for timely preventive interventions to reduce the impact of HZO** and its associated **healthcare burden**.

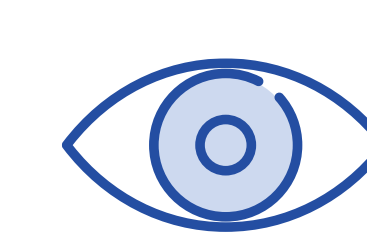
## Results

### Baseline demographics (N=104)

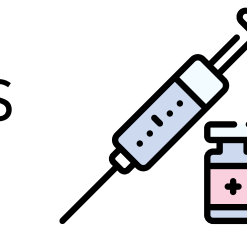
59.6% (n=62) female



Mean: **64.3 years** (SD: 13.7)



**70.2% (n=73)** initial HZO cases with **29.8% (n=31)** recurrent



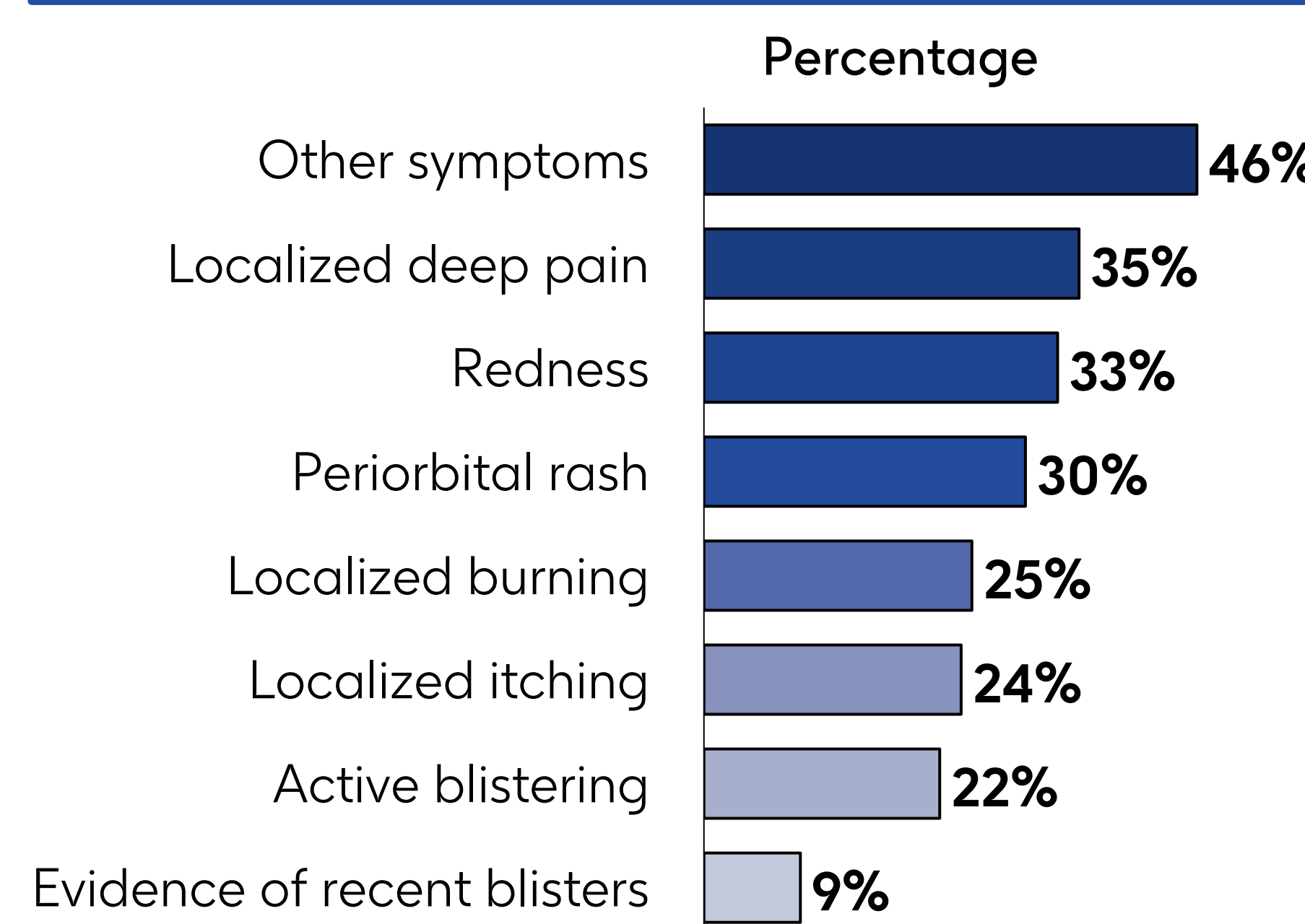
**28.8% (n=30)** vaccinated against herpes zoster



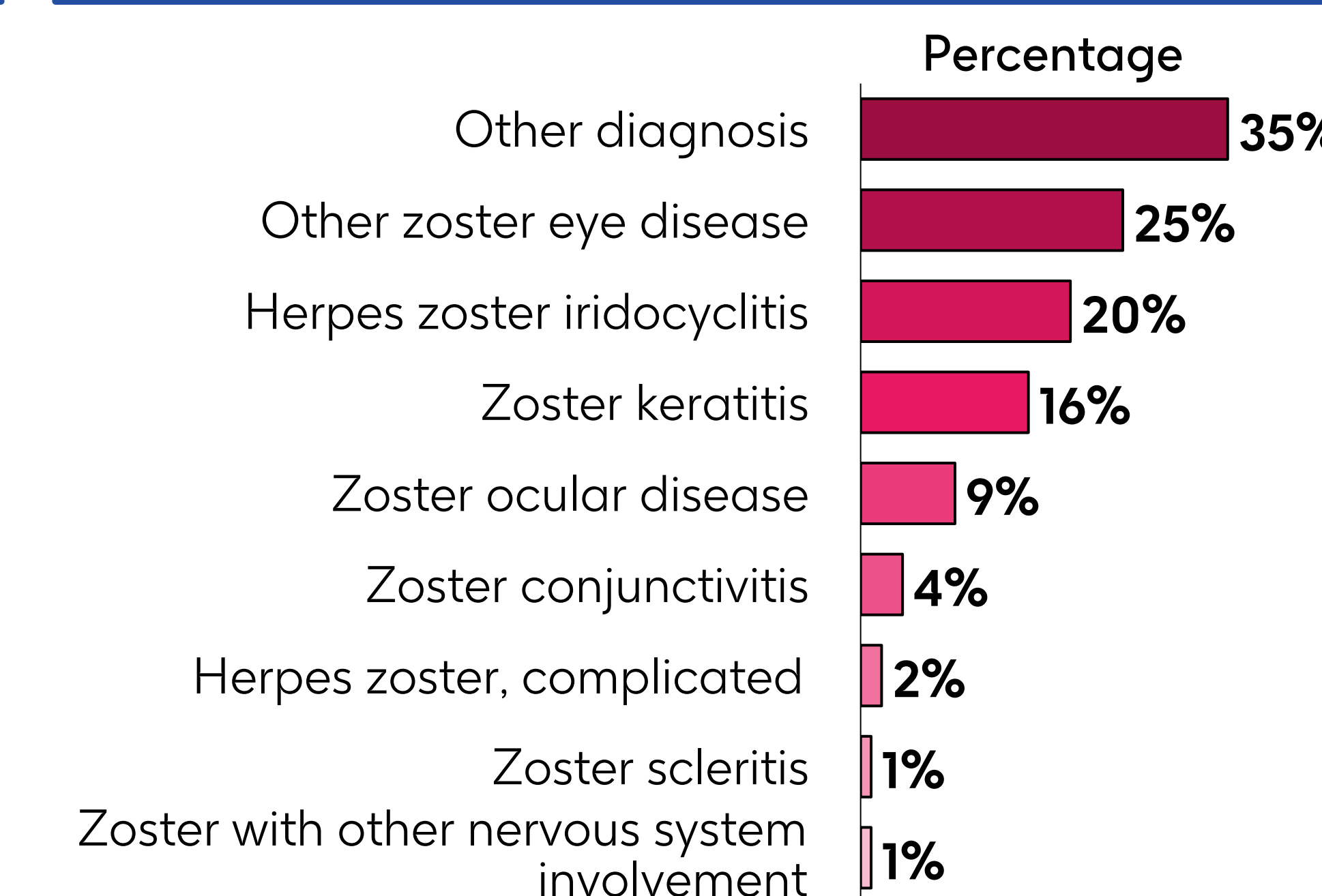
**66.4% (n=69)** White

### Baseline clinical characteristics, healthcare resource utilization, and patient-reported outcomes

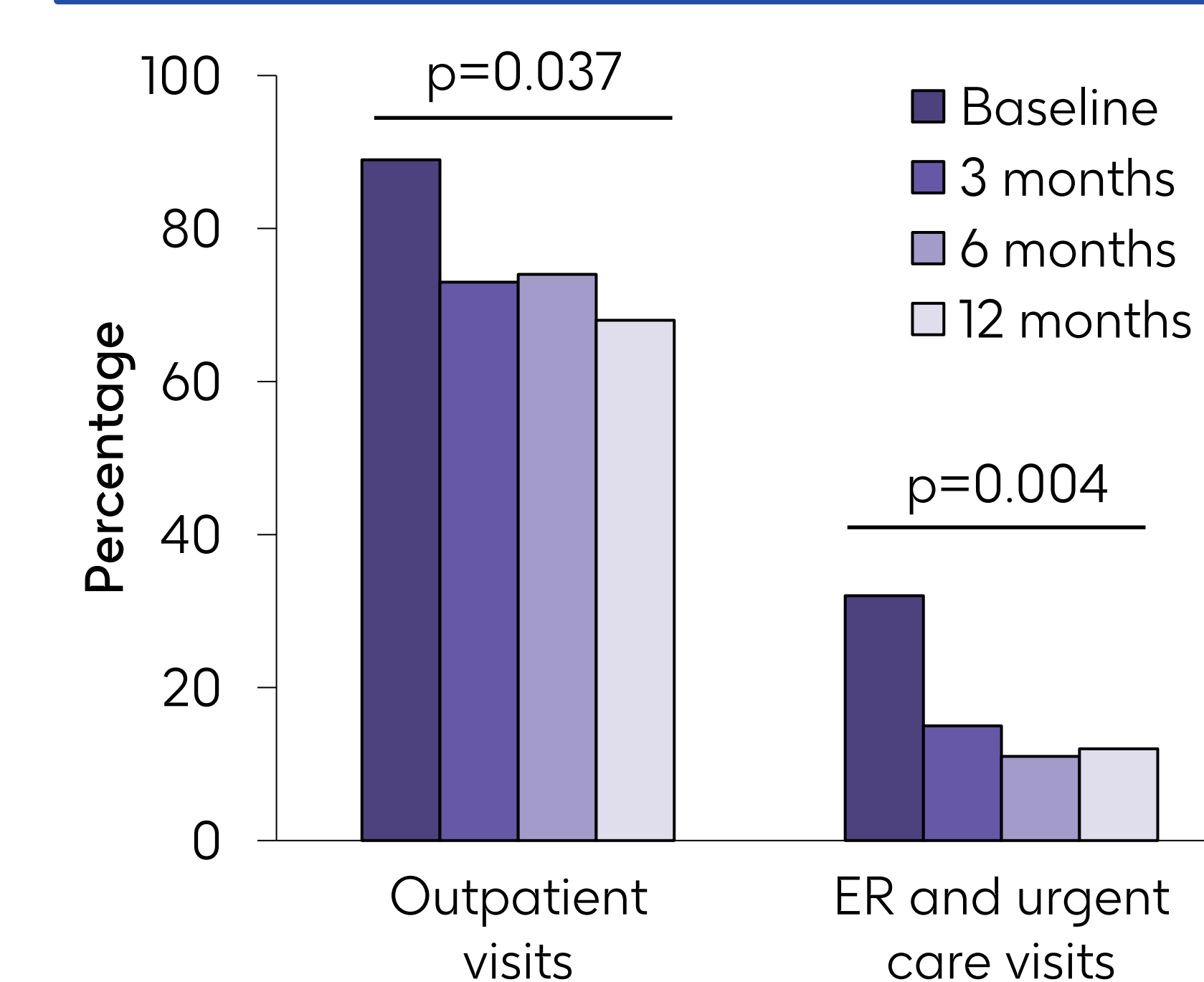
#### Clinical symptoms (N=110)<sup>a</sup>



#### Clinical diagnosis (N=110)<sup>a</sup>

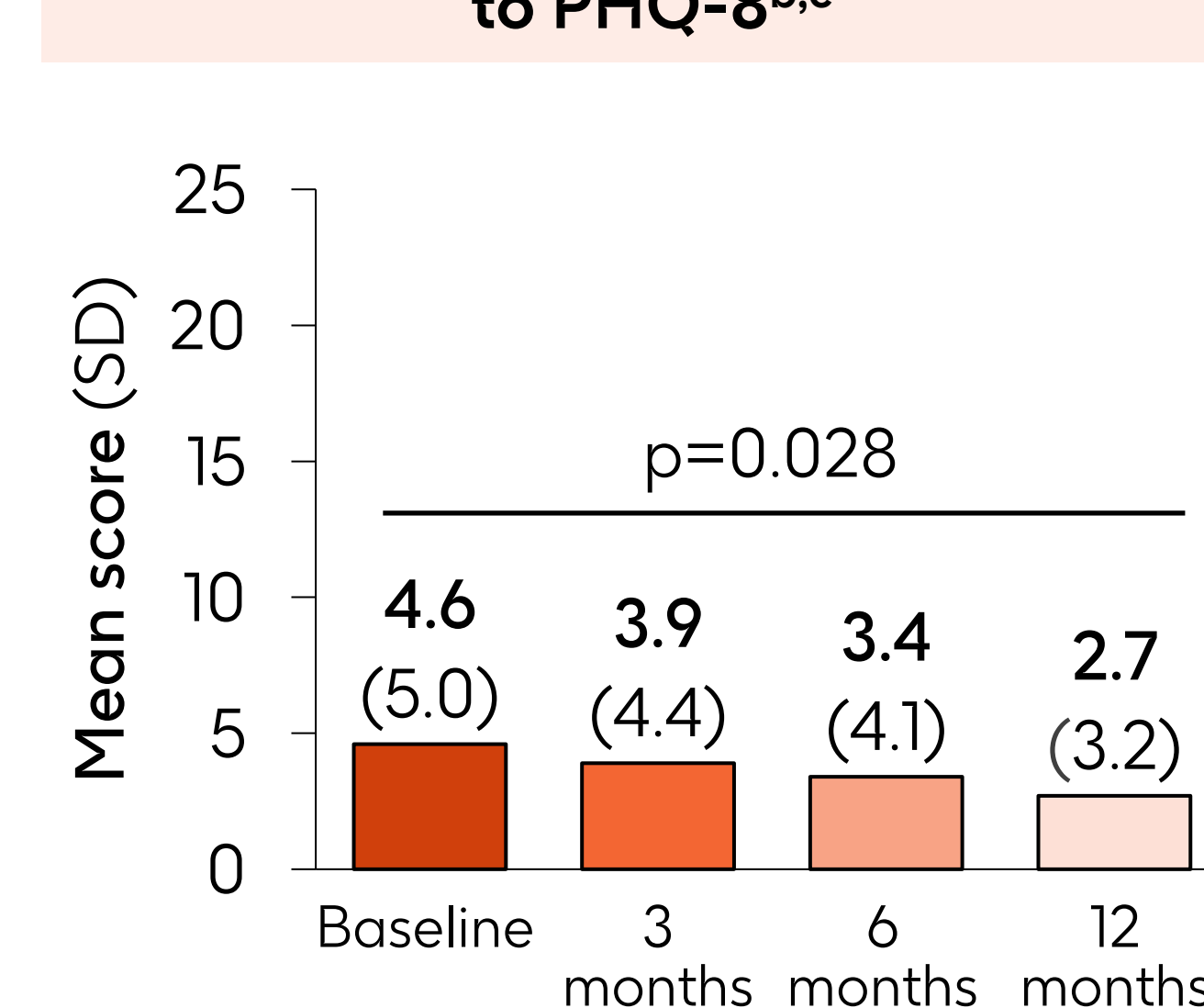


#### Healthcare resource utilization<sup>b</sup>

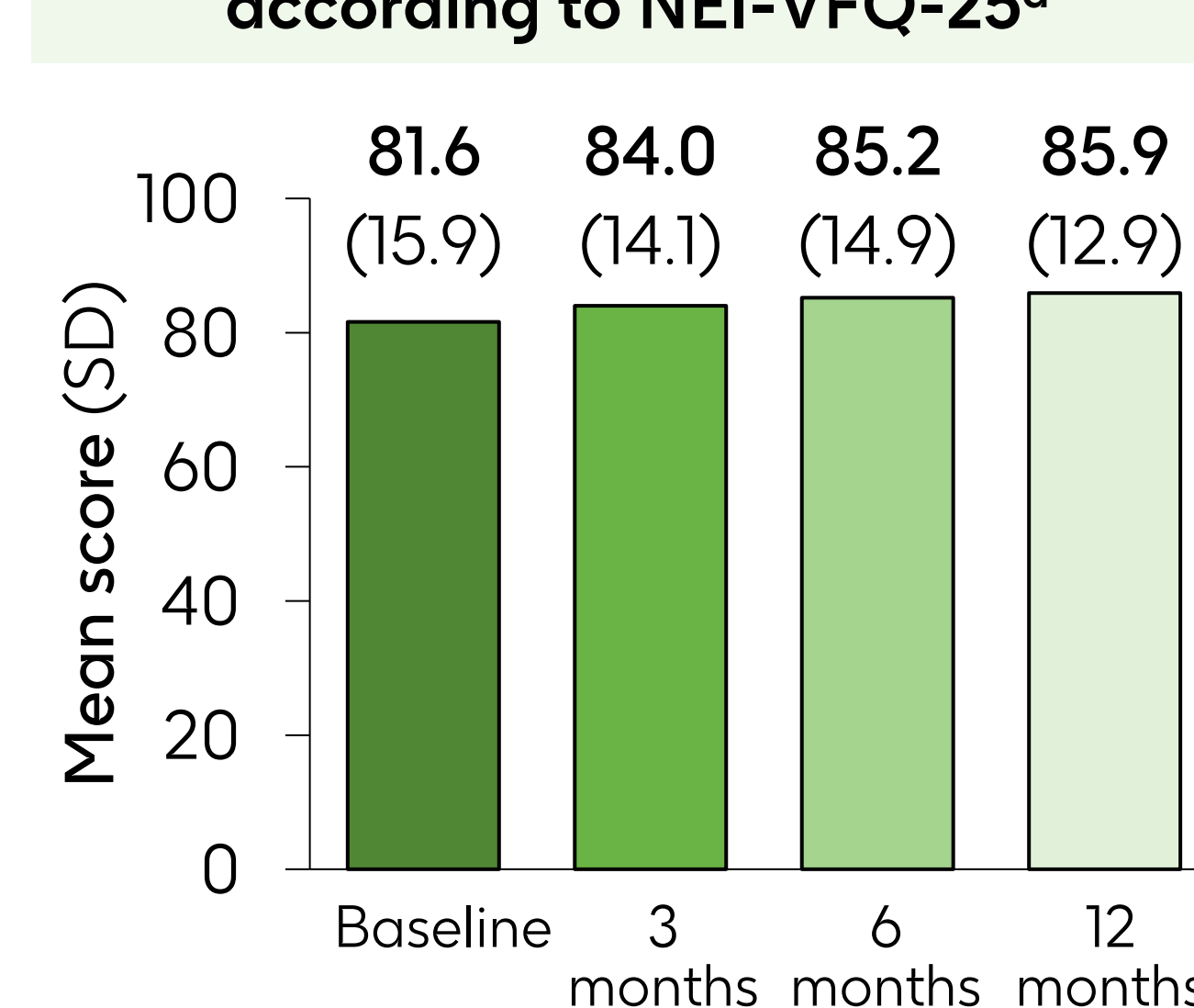


#### Patient-reported outcomes<sup>b</sup>

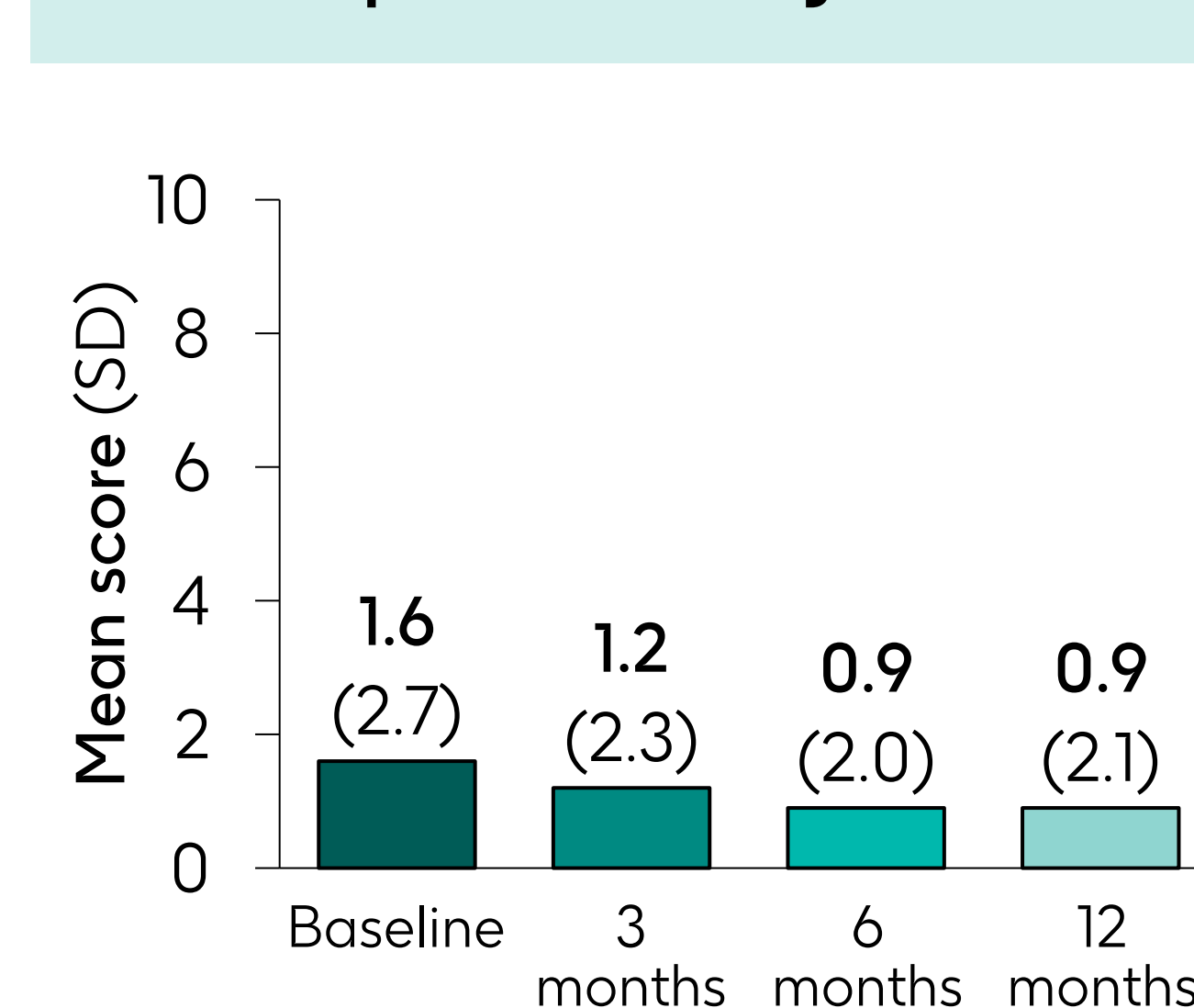
##### Depressive symptoms according to PHQ-8<sup>b,c</sup>



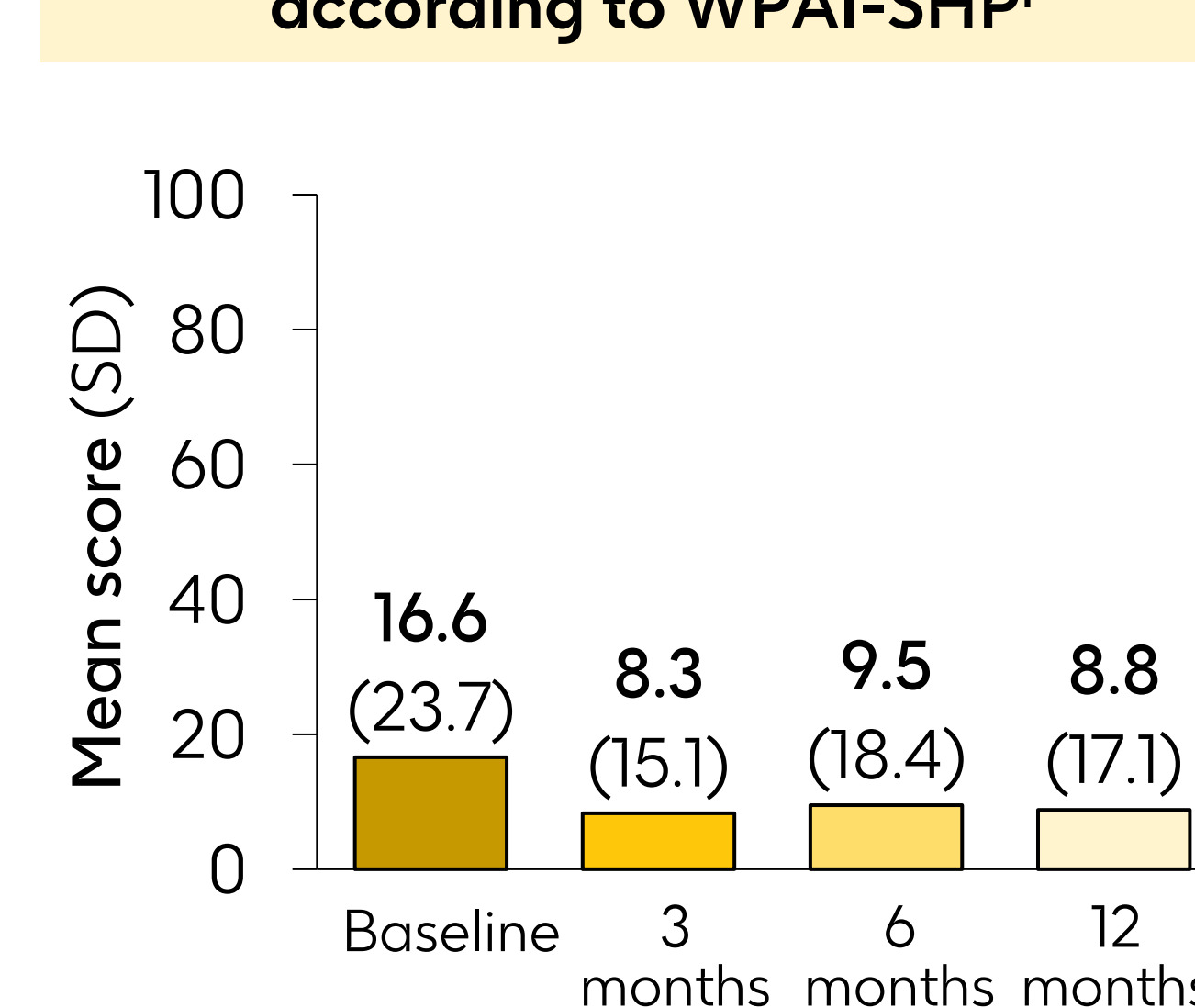
##### Vision-related quality of life according to NEI-VFQ-25<sup>d</sup>



##### Worst pain according to ZBPI<sup>e</sup>



##### Work productivity loss according to WPAI-SHP<sup>f</sup>



<sup>a</sup>104 patients completed the baseline survey while 110 patients completed at least one baseline or follow-up surveys (3, 6, or 12 months), the clinical characteristics are provided for 110 patients. See supplementary material (QR code) for additional details and data on clinical symptoms and diagnosis. <sup>b</sup>P-values are type-III p-values from multivariable regression analysis representing a significant difference between any of the time points. <sup>c</sup>Scores range from 0 to 24. Scores of 5, 10, 15, and 20 represent cutpoints for mild, moderate, moderately severe, and severe depression, respectively. <sup>d</sup>Scores range from 0 to 100. A high score represents better functioning. The scores should be interpreted as an achieved percentage of total possible score. <sup>e</sup>Scores range from 0 to 10. A high score represents worse pain severity from 0=no pain to 10=pain as bad as you can imagine. <sup>f</sup>Scores range from 0 to 100. Scores are expressed as impairment percentages, with higher numbers indicating greater impairment and less productivity.

## Abbreviations

ER: emergency room; HZO: herpes zoster ophthalmicus; N: total number of patients; n: number of patients; NEI-VFQ-25: National Eye Institute 25-Item Visual Function Questionnaire; PHQ-8: eight-item Patient Health Questionnaire; SD: standard deviation; US: United States; WPAI-SHP: Work Productivity and Activity Impairment-Specific Health Problem questionnaire; ZBPI: Zoster Brief Pain Inventory.

## References

- Giannelos N, et al. Infect Dis Ther. 2024;13(7):1461-86. 2. Dmitriev AA, et al. Can J Ophthalmol. 2024;59(3):201-7. 3. Yawn BP, et al. Mayo Clin Proc. 2013;88(6):562-70. 4. Curran D, et al. J Geriatr A Biol Sci Med Sci. 2019;74(8):1231-8. 5. Katz J, et al. Clin Infect Dis. 2004;39(3):342-8. 6. Kroenke K, et al. J Affect Disord. 2009;114(1-3):163-73. 7. Mangione CM, et al. Arch Ophthalmol. 1998;116(11):1496-504. 8. Mangione CM, et al. Arch Ophthalmol. 2001;119(7):1050-8. 9. Coplan PM, et al. J Pain. 2004;5(6):344-56. 10. Reilly et al. Pharmacoeconomics. 1993;4(5):353-65.

## Acknowledgments

Enovall Medical Communication Service Center provided editorial assistance and publications coordination, on behalf of GSK. The authors would like to thank Ms. Juliet Pettito (Rutgers University Ernest Mario School of Pharmacy) for her contribution to the study, Dr. David S. Chu (Metropolitan Eye Research and Surgery Institute / Rutgers New Jersey Medical School) participated in the data acquisition and in the interpretation of the study, and in the development and approval of the abstract.

## Disclosures

**Funding:** GSK (GSK study identifiers: 209235/HO-17-17967).

**Conflicts of interest:** LTP declares receipt of research funding from GSK for the submitted work. LTP also declares receipt of investigator-initiated research grant on vaccine hesitancy from Merck & Co., Inc., outside the submitted work, as well as being Associate Chief Science Officer of ISPOR—The Professional Society for Health Economics and Outcomes Research. BEL and AS declare their institution received a research grant from GSK for the submitted work. JPS declares contract from Rutgers University for patient recruitment for the present study. JPS also declares receipt of consulting fees from Bausch & Lomb; speaker honorarium from American Optometric Association, American Academy of Ophthalmology and American Academy of Optometry; payment for expert testimony from Willingham & Cote, Walsh, Barnes & Zumpella and Mongello Law; and support for attending meetings and/or travel from Optometric Management, outside the submitted work. BRF declares planned receipt of support for meeting attendance from UNC Chapel Hill Department of Ophthalmology outside the submitted work. KMP declares receipt of funding from GSK to her employer for the submitted work. KB and AK are employed by GSK and contracted through Center of Health Outcomes, Policy and Economics at Rutgers University. JDG is employed by, and holds financial equities in, GSK. JDG declares grants to his institution from Merck & Co., AstraZeneca, and GSK; payment or honoraria for lectures, presentations, speakers' bureaus, manuscript writing, or educational events from Merck & Co.; and reimbursement for attending meetings and/or travel from Genentech, outside the submitted work. APM declares receipt of grants and consulting fees from GSK for the submitted work. LTP, BEL, AS, JPS, BRF, KMP, KB, AK, JDG, and APM declare no other financial and non-financial relationships and activities. HJL, EWG, and HA declare no financial and non-financial relationships and activities, and no conflicts of interest.

