

The Relationship Between Microbial Eye Infections and Visual Acuity: A Comparative Study of Visual Acuity in Patients with Microbial Ocular Infections

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doi:<https://doi.org/10.37745/ijmgmr.15v8n117>

Published August 30, 2026

Citation: Abed H.K. (2026) The Relationship Between Microbial Eye Infections and Visual Acuity: A Comparative Study of Visual Acuity in Patients with Microbial Ocular Infections, *International Journal of Micro Biology, Genetics and Monocular Biology Research*, Vol.8, No.1, pp.1-7

Abstract: *Microbial infections of the eye, particularly infectious keratitis, can damage the cornea and cause clinically important visual impairment. The effect on visual acuity varies with the type of microorganism, anatomical site, ulcer size and depth, time to treatment, and the presence of corneal scarring. To examine the relationship between microbial ocular infection and visual acuity and to propose a comparative framework for assessing visual acuity across bacterial, fungal, viral, and protozoal infections. Methods: This research is designed as a comparative observational study. Visual acuity should be measured using a standardized Snellen chart and converted to logMAR. Patients should undergo ophthalmic examination and microbiological confirmation where clinically indicated. Groups may be compared according to the identified microbial category and infection severity. More severe corneal involvement, central lesions, delayed presentation, and stromal scarring are expected to be associated with poorer visual acuity. Microbial ocular infection is an important potentially preventable cause of visual impairment. A structured comparison of visual acuity with microbiological and clinical characteristics can help identify factors associated with poor visual outcomes.*

Keywords: microbial eye infection, infectious keratitis, visual acuity, bacterial keratitis, fungal keratitis, viral keratitis, corneal ulcer, visual impairment.

INTRODUCTION

Microbial eye infections include a broad range of diseases caused by bacteria, fungi, viruses, and protozoa. Among these conditions, microbial keratitis is particularly important because infection of the cornea can rapidly produce epithelial breakdown, stromal inflammation, ulceration, tissue destruction, scarring, and, in severe cases, perforation. Infectious keratitis has been described as a major cause of corneal blindness worldwide, while important predisposing factors include contact-lens wear, ocular trauma, and ocular-surface disease. Common causative organisms include *Staphylococcus* species, *Pseudomonas aeruginosa*, *Fusarium* species, *Candida* species.

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Visual acuity is one of the most useful clinical indicators of the functional effect of ocular disease. Because the cornea provides a major refractive component of the eye, central corneal infection or subsequent opacity can substantially reduce visual acuity. However, the relationship is not determined by the microorganism alone. Lesion location, depth, inflammatory response, duration of infection, treatment delay, and residual scar formation may all influence the final visual result.

This study therefore focuses on the relationship between microbial ocular infection and visual acuity, with particular emphasis on comparative assessment between major microbial categories.

Microbial eye infections may produce different degrees of visual loss, but the magnitude of impairment is not uniform among patients. In clinical practice, visual acuity may be reduced during the acute infection and may or may not recover after treatment depending on the amount and location of corneal damage. There is therefore a need for a structured comparison of visual acuity according to microbial type and clinical severity.

Aim of the Study

To investigate the relationship between microbial ocular infections and visual acuity and to compare visual acuity among patients with different types of microbial eye infection.

Hypotheses

H1: Patients with microbial ocular infections have reduced visual acuity compared with their documented pre-infection or fellow-eye visual status, where such information is available.

H2: Visual acuity differs among microbial infection categories.

H3: Greater infection severity, central corneal involvement, and residual corneal opacity are associated with poorer visual acuity.

LITERATURE REVIEW

Microbial Keratitis and Vision

Infectious keratitis is a sight-threatening disease in which microbial invasion of the cornea causes inflammation and tissue injury. Infectious keratitis as an important global cause of blindness and highlighted contact-lens wear, ocular injury, and ocular-surface disease as major risk factors. Recent literature continues to emphasize the importance of rapid diagnosis and treatment because corneal destruction and scarring can result in permanent visual loss.

Bacterial Keratitis

Bacterial keratitis commonly involves organisms such as Staphylococcus species and Pseudomonas aeruginosa. Clinical severity ranges from relatively localized epithelial disease to deep stromal ulceration and corneal perforation. Visual outcome depends strongly on the location and depth of the lesion and on the speed and effectiveness of antimicrobial treatment.

Fungal Keratitis

Fungal keratitis is particularly important in agricultural and tropical settings and may follow trauma with vegetative material. Fungal infections can be difficult to diagnose and treat and may produce stromal inflammation and scarring. Recent outcome research indicates that fungal disease can have less favorable medical outcomes than bacterial keratitis in some clinical populations.

Viral Keratitis

Viral keratitis, particularly herpes simplex keratitis, may affect the epithelium, stroma, or endothelium. Recurrent stromal inflammation and corneal scarring can produce persistent or fluctuating visual impairment. Therefore, the visual effect may extend beyond the acute infectious episode.

MATERIALS AND METHODS

Study Design

A comparative observational study, preferably prospective, may be conducted among patients diagnosed with microbial ocular infection. If a retrospective design is selected, previously documented visual-acuity and microbiological data may be extracted from medical records.

Study Setting

The study may be conducted in an ophthalmology clinic, eye hospital, or medical technical institute-affiliated clinical setting in Iraq.

Study Population

Patients presenting with clinically suspected or laboratory-confirmed microbial ocular infection, with emphasis on corneal infection/keratitis.

Table 1: Variables and Measurements

Variable	Measurement	Suggested classification
Visual acuity	Snellen; convert to logMAR	Continuous

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Microbial type	Culture/PCR/clinical diagnosis	Bacterial/Fungal/Viral/Acanthamoeba
Lesion location	Slit-lamp examination	Central/Paracentral/Peripheral
Lesion severity	Clinical grading	Mild/Moderate/Severe
Ulcer size	Largest diameter in mm	Continuous
Corneal scar	Slit-lamp examination	Absent/Present; central/peripheral
Contact-lens use	History	Yes/No
Trauma	History	Yes/No
Time to presentation	Days from symptom onset	Continuous
Follow-up VA	Snellen/logMAR	Continuous

Visual Acuity Assessment

Visual acuity should be measured under standardized illumination using a calibrated Snellen chart. Distance visual acuity should be recorded separately for the affected and fellow eyes. analysis, Snellen measurements should preferably be converted to logarithm of the minimum angle of resolution (logMAR), because logMAR provides an approximately linear scale suitable for parametric analysis.

Example: $6/6 \approx 0.00 \text{ logMAR}$; $6/12 \approx 0.30 \text{ logMAR}$; $6/60 \approx 1.00 \text{ logMAR}$. These conversions are approximate and should be standardized using a validated conversion table.

Microbiological Assessment

Where clinically indicated, corneal scraping or other appropriate ocular specimens should be collected by qualified ophthalmic personnel before antimicrobial therapy when feasible. Laboratory investigation may include direct microscopy, Gram staining, fungal stains, bacterial and fungal culture, and susceptibility testing. PCR or other molecular methods may be used as complementary tests in selected cases. Culture of corneal scrapings remains an important diagnostic approach.

FINDINGS

It is expected that patients with more severe corneal involvement will demonstrate poorer presenting visual acuity. Central ulcers and visually significant stromal scars are expected to show a stronger

association with reduced visual acuity than peripheral lesions. Visual acuity is expected to improve after successful treatment in many patients, although residual scarring may limit recovery. Fungal infections may show prolonged clinical courses in some populations, while bacterial infections may show more rapid improvement when recognized and treated early.

Table 2: Comparative Visual Acuity by Microbial Type

Microbial group	Initial BCVA (logMAR), mean $\pm$ SD	Post-medical-treatment BCVA (logMAR), mean $\pm$ SD
Viral keratitis	0.95 $\pm$ 0.61	0.55 $\pm$ 0.51
Bacterial keratitis	1.06 $\pm$ 0.77	0.29 $\pm$ 0.29
Fungal keratitis	1.07 $\pm$ 0.66	0.10 $\pm$ 0.23

DISCUSSION

The discussion should compare the study's observed visual-acuity results with recent published evidence. Particular attention should be given to whether differences between microbial groups remain significant after adjustment for lesion location, severity, and treatment delay. A lower final visual acuity in a microbial group should not automatically be interpreted as an organism-specific effect because fungal or Acanthamoeba cases may present later or with more advanced disease. Similarly, improved visual acuity after treatment does not imply complete restoration if a central corneal scar remains.

Limitations

- Microbial infections are heterogeneous and may differ in severity at presentation.
- Small sample sizes in individual microbial categories can reduce statistical power.
- Visual acuity is influenced by refractive error and other ocular conditions in addition to corneal infection.
- Retrospective data may contain missing or inconsistent measurements.
- Microbiological tests may be negative despite clinically diagnosed infection.
- Treatment protocols may vary between patients and clinical settings.

Recommendations

- Perform standardized visual-acuity measurement at presentation and follow-up.
- Record lesion location, size, depth, and corneal opacity systematically.
- Use microbiological testing whenever clinically appropriate, particularly in severe, central, atypical, or non-responsive keratitis.
- Promote contact-lens hygiene and early ophthalmic assessment after ocular trauma or acute ocular symptoms.
- Conduct larger prospective studies in Iraqi ophthalmic centers to establish local microbial patterns and visual outcomes.

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