

**K772221 LENS, HARD RESIN**Jan 30, 1978  
56 days to decisionK772221 · Product code: **HQG** · Ophthalmic  
Source: <https://www.510kdatabase.net/k772221/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lens, Spectacle, Non-custom (prescription) (HQG)
Date received	Dec 5, 1977
Decision date	Jan 30, 1978
Days to decision	56 days
Third-party review	No

**APPLICANT**

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Company	<b>Borani Intl.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1978-1978

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k772221/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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