

**K782141 LEVIGATOR, KEELER PHOKO**Jan 26, 1979  
30 days to decisionK782141 · Product code: **HQC** · Ophthalmic  
Source: <https://www.510kdatabase.net/k782141/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Phacofragmentation (HQC)
Date received	Dec 27, 1978
Decision date	Jan 26, 1979
Days to decision	30 days
Third-party review	No

**APPLICANT**

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Company	<b>Keeler Optical Products , Ltd.</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1978-1981

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Device record: <https://www.510kdatabase.net/k782141/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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