FDA 510(k) Premarket Notification

K782141 LEVIGATOR, KEELER PHOKO

CLEARED

Jan 26, 1979
30 days to decision

K782141 · Product code: **HQC** · Ophthalmic
Source: https://www.510kdatabase.net/k782141/

SUBMISSION DETAILS

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

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