FDA 510(k) Premarket Notification

K801969 ASNIS GUIDED SCREW SYSTEMS

CLEARED

Aug 27, 1980
8 days to decision

K801969 · Product code: **HNC** · Ophthalmic
Source: https://www.510kdatabase.net/k801969/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Specula, Ophthalmic (HNC)
Date received	Aug 19, 1980
Decision date	Aug 27, 1980
Days to decision	8 days
Third-party review	No

APPLICANT

Company	**Howmedica Corp.**
Location	Mchenry, IL, US
510(k) history	373 submissions · 325 cleared · 1976-1998