

K801969 ASNIS GUIDED SCREW SYSTEMSAug 27, 1980
8 days to decisionK801969 · 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

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Device record: <https://www.510kdatabase.net/k801969/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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