

K830430 EYELID SPECULUMJun 10, 1983
121 days to decisionK830430 · Product code: **HNC** · Ophthalmic
Source: <https://www.510kdatabase.net/k830430/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Specula, Ophthalmic (HNC)
Date received	Feb 9, 1983
Decision date	Jun 10, 1983
Days to decision	121 days
Third-party review	No

APPLICANT

Company	Douglas C. Mckee & Co.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1983-1983

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k830430/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 1, 2026