FDA 510(k) Premarket Notification

K830430 EYELID SPECULUM

CLEARED

Jun 10, 1983
121 days to decision

K830430 · 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