FDA 510(k) Premarket Notification

K811455 ANIS GUIDING CANNULA

CLEARED

Jun 16, 1981
25 days to decision

K811455 · Product code: **HMX** · Ophthalmic
Source: https://www.510kdatabase.net/k811455/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Ophthalmic (HMX)
Date received	May 22, 1981
Decision date	Jun 16, 1981
Days to decision	25 days
Third-party review	No

APPLICANT

Company	**American V. Mueller**
Location	Mchenry, IL, US
510(k) history	31 submissions · 29 cleared · 1980-1987