

K811455 ANIS GUIDING CANNULAJun 16, 1981
25 days to decisionK811455 · 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

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Device record: <https://www.510kdatabase.net/k811455/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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