

K782169 NEEDLES, CANNULAS, CYSTOTOMESFeb 13, 1979
46 days to decisionK782169 · Product code: **HMX** · Ophthalmic
Source: <https://www.510kdatabase.net/k782169/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Ophthalmic (HMX)
Date received	Dec 29, 1978
Decision date	Feb 13, 1979
Days to decision	46 days
Third-party review	No

APPLICANT

Company	Visitec Co.
Location	Walker, MI, US
510(k) history	49 submissions · 49 cleared · 1979-1995

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k782169/> 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 May 23, 2026