

**K790243 CANNULAS**Feb 12, 1979  
45 days to decisionK790243 · Product code: **HMX** · Ophthalmic  
Source: <https://www.510kdatabase.net/k790243/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Ophthalmic (HMX)
Date received	Dec 29, 1978
Decision date	Feb 12, 1979
Days to decision	45 days
Third-party review	No

**APPLICANT**

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

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Device record: <https://www.510kdatabase.net/k790243/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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