

K831901 LACIMAL INTUBATION SETSJul 12, 1983
28 days to decisionK831901 · Product code: **HNL** · Ophthalmic
Source: <https://www.510kdatabase.net/k831901/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Lachrymal (HNL)
Date received	Jun 14, 1983
Decision date	Jul 12, 1983
Days to decision	28 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k831901/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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