FDA 510(k) Premarket Notification

K831901 LACIMAL INTUBATION SETS

CLEARED

Jul 12, 1983
28 days to decision

K831901 · 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