FDA 510(k) Premarket Notification

K902857 VISI-DRAPE OPHTHALMIC DRAPES

CLEARED

Jul 31, 1990
32 days to decision

K902857 · Product code: **HMY** · Ophthalmic
Source: https://www.510kdatabase.net/k902857/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Keratome, Battery-powered (HMY)
Date received	Jun 29, 1990
Decision date	Jul 31, 1990
Days to decision	32 days
Third-party review	No

APPLICANT

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