FDA 510(k) Premarket Notification

K884924 VISI-WIPE

CLEARED

Jan 31, 1989
64 days to decision

K884924 · Product code: **HOF** · Ophthalmic
Source: https://www.510kdatabase.net/k884924/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Burr, Corneal, Manual (HOF)
Date received	Nov 28, 1988
Decision date	Jan 31, 1989
Days to decision	64 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