FDA 510(k) Premarket Notification

K960395 PLANCON MIRCOLAMELLAR KERATOME

CLEARED

Apr 26, 1996
88 days to decision

K960395 · Product code: **HNO** · Ophthalmic
Source: https://www.510kdatabase.net/k960395/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Keratome, Ac-powered (HNO)
Date received	Jan 29, 1996
Decision date	Apr 26, 1996
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Plancon Instruments**
Location	Antony, FR
Contact	ALAIN DUPRAT
510(k) history	3 submissions · 3 cleared · 1996-1998