FDA 510(k) Premarket Notification

K903912 KERATOME

CLEARED

Oct 17, 1990
54 days to decision

K903912 · Product code: **HNO** · Ophthalmic
Source: https://www.510kdatabase.net/k903912/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Keratome, Ac-powered (HNO)
Date received	Aug 24, 1990
Decision date	Oct 17, 1990
Days to decision	54 days
Third-party review	No

APPLICANT

Company	**Micro Precision Instrument Co.**
Location	Scottsdale, AR, US
Contact	RUSSELL KOEPNICK
510(k) history	1 submissions · 1 cleared · 1990-1990