FDA 510(k) Premarket Notification

K923708 LEMPERT OPTIC DISC ANALYSIS SYSTEM

CLEARED

Mar 19, 1993
235 days to decision

K923708 · Product code: **HKI** · Ophthalmic
Source: https://www.510kdatabase.net/k923708/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Ophthalmic, Ac-powered (HKI)
Date received	Jul 27, 1992
Decision date	Mar 19, 1993
Days to decision	235 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Lempert Ophthalmic Instruments**
Location	Ithaca, NY, US
Contact	PHILIP LEMPERT
510(k) history	1 submissions · 1 cleared · 1993-1993