FDA 510(k) Premarket Notification

K912740 ACT-1 (ANTERIOR CORNEAL TOPOGRAPHER)

CLEARED

Dec 30, 1991
192 days to decision

K912740 · Product code: **HLQ** · Ophthalmic
Source: https://www.510kdatabase.net/k912740/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Keratoscope, Ac-powered (HLQ)
Date received	Jun 21, 1991
Decision date	Dec 30, 1991
Days to decision	192 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Visionary Systems, Inc.**
Location	Tucson, AZ, US
Contact	STEVEN R LANGE
510(k) history	1 submissions · 1 cleared · 1991-1991