

K912740 ACT-1 (ANTERIOR CORNEAL TOPOGRAPHER)Dec 30, 1991
192 days to decisionK912740 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k912740/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 30, 2026