FDA 510(k) Premarket Notification

K972252 VISTA UNIVERSAL HANDPIECE

CLEARED

Jul 29, 1997
43 days to decision

K972252 · Product code: **HQC** · Ophthalmic
Source: https://www.510kdatabase.net/k972252/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Phacofragmentation (HQC)
Date received	Jun 16, 1997
Decision date	Jul 29, 1997
Days to decision	43 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Odyssey Technologies, Inc.**
Location	Temple City, CA, US
Contact	GLENN A DUNKI-JACOBS
510(k) history	2 submissions · 2 cleared · 1997-1997