FDA 510(k) Premarket Notification

K071973 EPLEY OMNIAX

CLEARED

Jun 20, 2008
339 days to decision

K071973 · Product code: **LXV** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k071973/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Vestibular Analysis (LXV)
Date received	Jul 17, 2007
Decision date	Jun 20, 2008
Days to decision	339 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Vesticon**
Location	Portland, OR, US
Contact	CATHRYN EPLEY
510(k) history	1 submissions · 1 cleared · 2008-2008