FDA 510(k) Premarket Notification

K904909 I/A VITROPHAGE MODEL YPR 2001

CLEARED

Jan 28, 1991
89 days to decision

K904909 · Product code: **HQE** · Ophthalmic
Source: https://www.510kdatabase.net/k904909/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Vitreous Aspiration And Cutting, Ac-powered (HQE)
Date received	Oct 31, 1990
Decision date	Jan 28, 1991
Days to decision	89 days
Third-party review	No

APPLICANT

Company	**Visioncare Devices, Inc.**
Location	Redding, CA, US
Contact	CHET CRACCHIOLO
510(k) history	5 submissions · 5 cleared · 1991-2012