FDA 510(k) Premarket Notification

K894248 CONVERTER SYSTEM

CLEARED

Feb 27, 1990
251 days to decision

K894248 · Product code: **HQE** · Ophthalmic
Source: https://www.510kdatabase.net/k894248/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Vitreous Aspiration And Cutting, Ac-powered (HQE)
Date received	Jun 21, 1989
Decision date	Feb 27, 1990
Days to decision	251 days
Third-party review	No

APPLICANT

Company	**Mira, Inc.**
Location	Waltham, MA, US
Contact	ROGER O'BRIEN
510(k) history	21 submissions · 21 cleared · 1984-1997