FDA 510(k) Premarket Notification

K020294 MOTION TRACKING

CLEARED

Apr 22, 2002
84 days to decision

K020294 · Product code: **IYE** · Radiology
Source: https://www.510kdatabase.net/k020294/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Jan 28, 2002
Decision date	Apr 22, 2002
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Accuray, Inc.**
Location	Sunnyvale, CA, US
Contact	DONALD E CADDES
510(k) history	10 submissions · 10 cleared · 1999-2009