FDA 510(k) Premarket Notification

K902334 MONITOR, MODEL 2817

CLEARED

Jun 26, 1990
33 days to decision

K902334 · Product code: **DXJ** · Cardiovascular
Source: https://www.510kdatabase.net/k902334/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Display, Cathode-ray Tube, Medical (DXJ)
Date received	May 24, 1990
Decision date	Jun 26, 1990
Days to decision	33 days
Third-party review	No

APPLICANT

Company	**Esaote Biomedica Spa**
Location	Cambridge, MA, US
Contact	BARRY SALL
510(k) history	4 submissions · 4 cleared · 1990-1994