FDA 510(k) Premarket Notification

K811911 SINGLE CHANNEL NON-FADE OSCILLOSCOPE

CLEARED

Jul 28, 1981
27 days to decision

K811911 · Product code: **DXJ** · Cardiovascular
Source: https://www.510kdatabase.net/k811911/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Display, Cathode-ray Tube, Medical (DXJ)
Date received	Jul 1, 1981
Decision date	Jul 28, 1981
Days to decision	27 days
Third-party review	No

APPLICANT

Company	**Quinton, Inc.**
Location	Mchenry, IL, US
510(k) history	164 submissions · 160 cleared · 1976-2003