

K811911 SINGLE CHANNEL NON-FADE OSCILLOSCOPEJul 28, 1981
27 days to decisionK811911 · 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

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Device record: <https://www.510kdatabase.net/k811911/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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