FDA 510(k) Premarket Notification

K810563 QUINTON LTE ELECTRODE

CLEARED

Mar 17, 1981
14 days to decision

K810563 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k810563/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Mar 3, 1981
Decision date	Mar 17, 1981
Days to decision	14 days
Third-party review	No

APPLICANT

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