FDA 510(k) Premarket Notification

K782079 ELECTRODE SYSTEM, QUIK-PREP

CLEARED

Feb 1, 1979
50 days to decision

K782079 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k782079/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Dec 13, 1978
Decision date	Feb 1, 1979
Days to decision	50 days
Third-party review	No

APPLICANT

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