FDA 510(k) Premarket Notification

K760280 ELECTRODE SERIES 1401

CLEARED

Aug 23, 1976
31 days to decision

K760280 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k760280/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jul 23, 1976
Decision date	Aug 23, 1976
Days to decision	31 days
Third-party review	No

APPLICANT

Company	**Andover Medical, Inc.**
Location	Mchenry, IL, US
Website	https://www.andovermedical.com
510(k) history	21 submissions · 21 cleared · 1976-1991