FDA 510(k) Premarket Notification

K991105 ECG ELECTRODE (VARIOUS)

CLEARED

Nov 26, 1999
239 days to decision

K991105 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k991105/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Apr 1, 1999
Decision date	Nov 26, 1999
Days to decision	239 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Conmedcorp**
Location	Dayton, OH, US
Contact	IRA D DUESLER
510(k) history	92 submissions · 92 cleared · 1981-2010