FDA 510(k) Premarket Notification

K954388 CONMED 2710 CLEARTRACE 2 LT MONITORING ECG ELECTRODE

CLEARED

Feb 6, 1996
140 days to decision

K954388 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k954388/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Sep 19, 1995
Decision date	Feb 6, 1996
Days to decision	140 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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