FDA 510(k) Premarket Notification

K091856 CONMED ECG MONITORING ELECTRODES , MODEL 1700 (INCLUDING 2500), 1750, 2700, 2710, 1800, SERIES

CLEARED

Aug 7, 2009
45 days to decision

K091856 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k091856/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jun 23, 2009
Decision date	Aug 7, 2009
Days to decision	45 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Conmed Corporation**
Location	Utica, NY, US
Contact	SANDRA COVELESKI
Website	https://www.conmed.com
510(k) history	83 submissions · 83 cleared · 2004-2026

Conmed Corporation is a global medical device manufacturer specializing in surgical equipment and operating room solutions. The company operates with a manufacturing facility in Utica, US, and serves multiple surgical specialties including general surgery, orthopedics, and patient monitoring. Conmed has received FDA 510(k) clearances from total submissions since its first clearance in 2004. The company maintains active regulatory engagement, with its most recent clearance in 2026. Its cleared devices focus primarily on General & Plastic Surgery applications, including ele...