FDA 510(k) Premarket Notification

K962122 SENTRY RADIOTRANSLUCENT ECG ELECTRODES

CLEARED

Jul 24, 1996
54 days to decision

K962122 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k962122/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	May 31, 1996
Decision date	Jul 24, 1996
Days to decision	54 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Sentry Medical Products, Inc.**
Location	Walker, MI, US
Contact	JAMES G STROKES
510(k) history	13 submissions · 13 cleared · 1983-1996