FDA 510(k) Premarket Notification

K944260 ECG ELECTRODE

CLEARED

Jul 7, 1995
311 days to decision

K944260 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k944260/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Aug 30, 1994
Decision date	Jul 7, 1995
Days to decision	311 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Msb , Ltd.**
Location	Marlborough, Wiltshire, GB
Contact	MARTIN BEAUMONT
510(k) history	7 submissions · 7 cleared · 1994-1998