FDA 510(k) Premarket Notification

K002310 CARDIOBEEPER CB12/12, MODEL CB12/12

CLEARED

Oct 25, 2000
86 days to decision

K002310 · Product code: **DXH** · Cardiovascular
Source: https://www.510kdatabase.net/k002310/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Electrocardiograph, Telephone (DXH)
Date received	Jul 31, 2000
Decision date	Oct 25, 2000
Days to decision	86 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Meridian Medical Technologies , Ltd.**
Location	Belfast, IE
Contact	GERARD LYNN
510(k) history	3 submissions · 3 cleared · 2000-2001