

K002310 CARDIOBEEPER CB12/12, MODEL CB12/12Oct 25, 2000
86 days to decisionK002310 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k002310/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 28, 2026