FDA 510(k) Premarket Notification

K031210 MODEL 4951M MYOCARDIAL UNIPOLAR LEAD

CLEARED

May 16, 2003
29 days to decision

K031210 · Product code: **DTB** · Cardiovascular
Source: https://www.510kdatabase.net/k031210/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Apr 17, 2003
Decision date	May 16, 2003
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Metronic, Inc.**
Location	Mchenry, IL, US
Contact	TINA BENOIT
510(k) history	4 submissions · 4 cleared · 1983-2003