

K932254 MAXIMA FILTERED HARDSHELL RESERVOIRAug 3, 1993
85 days to decisionK932254 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k932254/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - SP
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	May 10, 1993
Decision date	Aug 3, 1993
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic Vascular
Location	Walker, MI, US
Contact	NORMA LOWE
510(k) history	475 submissions · 453 cleared · 1977-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k932254/> 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 May 14, 2026