

**K900515 MODELS 6207/6208/6209/6210/6211/6212/6214
(PLI'S)**Aug 23, 1990
202 days to decisionK900515 · Product code: **DYB** · Cardiovascular
Source: <https://www.510kdatabase.net/k900515/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Feb 2, 1990
Decision date	Aug 23, 1990
Days to decision	202 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k900515/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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