FDA 510(k) Premarket Notification

K900515 MODELS 6207/6208/6209/6210/6211/6212/6214 (PLI'S)

CLEARED

Aug 23, 1990
202 days to decision

K900515 · 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