FDA 510(k) Premarket Notification

K890898 MODEL 611101 CONTRAST INJECTION DEVICE

CLEARED

Apr 12, 1989
49 days to decision

K890898 · Product code: **DXT** · Cardiovascular
Source: https://www.510kdatabase.net/k890898/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Feb 22, 1989
Decision date	Apr 12, 1989
Days to decision	49 days
Third-party review	No

APPLICANT

Company	**Medtronic Versaflex, Inc.**
Location	San Diego, CA, US
Contact	TIMOTHY J JOHNSON
510(k) history	2 submissions · 2 cleared · 1989-1989