FDA 510(k) Premarket Notification

K830609 INFLATION DEVICE

CLEARED

Mar 24, 1983
27 days to decision

K830609 · Product code: **DXT** · Cardiovascular
Source: https://www.510kdatabase.net/k830609/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Feb 25, 1983
Decision date	Mar 24, 1983
Days to decision	27 days
Third-party review	No

APPLICANT

Company	**Interventional Medical, Inc.**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1982-1988