FDA 510(k) Premarket Notification

K904212 HIGH-PRESSURE CONNECTOR TUBE

CLEARED

Dec 3, 1990
82 days to decision

K904212 · Product code: **DXT** · Cardiovascular
Source: https://www.510kdatabase.net/k904212/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Sep 12, 1990
Decision date	Dec 3, 1990
Days to decision	82 days
Third-party review	No

APPLICANT

Company	**Mallinckrodt Medical**
Location	St Louis, MO, US
Contact	DAVID E BROWN
510(k) history	43 submissions · 42 cleared · 1990-2008