FDA 510(k) Premarket Notification

K823340 CLEAR VINYL MONITORING CONNECTOR

CLEARED

Dec 30, 1982
52 days to decision

K823340 · Product code: **DXT** · Cardiovascular
Source: https://www.510kdatabase.net/k823340/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Nov 8, 1982
Decision date	Dec 30, 1982
Days to decision	52 days
Third-party review	No

APPLICANT

Company	**Vertex Medical Corp.**
Location	Mchenry, IL, US
510(k) history	21 submissions · 21 cleared · 1982-1984