FDA 510(k) Premarket Notification

K823254 REUSABLE METAL STOPCOCK

CLEARED

Dec 30, 1982
59 days to decision

K823254 · Product code: **DXT** · Cardiovascular
Source: https://www.510kdatabase.net/k823254/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Nov 1, 1982
Decision date	Dec 30, 1982
Days to decision	59 days
Third-party review	No

APPLICANT

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