FDA 510(k) Premarket Notification

K823191 DISPOSABLE PLASTIC STOPCOCK

CLEARED

Dec 30, 1982
65 days to decision

K823191 · Product code: **DXT** · Cardiovascular
Source: https://www.510kdatabase.net/k823191/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Injector And Syringe, Angiographic (DXT)
Date received	Oct 26, 1982
Decision date	Dec 30, 1982
Days to decision	65 days
Third-party review	No

APPLICANT

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