FDA 510(k) Premarket Notification

K232536 Soldier Microcatheter

CLEARED

Feb 23, 2024
186 days to decision

K232536 · Product code: **DQO** · Cardiovascular
Source: https://www.510kdatabase.net/k232536/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Intravascular, Diagnostic (DQO)
Date received	Aug 21, 2023
Decision date	Feb 23, 2024
Days to decision	186 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Embolx, Inc.**
Location	Los Altos, CA, US
Contact	Louise Musante
510(k) history	3 submissions · 3 cleared · 2015-2024