

K232536 Soldier MicrocatheterFeb 23, 2024
186 days to decisionK232536 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k232536/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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