FDA 510(k) Premarket Notification

K181954 autoLog IQ Autotransfusion System

CLEARED

Aug 22, 2018
30 days to decision

K181954 · Product code: **CAC** · Cardiovascular
Source: https://www.510kdatabase.net/k181954/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Apparatus, Autotransfusion (CAC)
Date received	Jul 23, 2018
Decision date	Aug 22, 2018
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medtronic Perfusion Systems**
Location	Minneapolis, MN, US
Contact	Sue Fidler
510(k) history	29 submissions · 29 cleared · 2000-2024