

**K181954 autoLog IQ Autotransfusion System**Aug 22, 2018  
30 days to decisionK181954 · 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	<b>Medtronic Perfusion Systems</b>
Location	Minneapolis, MN, US
Contact	Sue Fidler
510(k) history	29 submissions · 29 cleared · 2000-2024

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k181954/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 14, 2026