

K150894 VIA 21 MicrocatheterAug 28, 2015
148 days to decisionK150894 · Product code: **DQY** · Cardiovascular
Source: <https://www.510kdatabase.net/k150894/>**SUBMISSION DETAILS**

| | |
|-----------------------|------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Percutaneous (DQY) |
| Date received | Apr 2, 2015 |
| Decision date | Aug 28, 2015 |
| Days to decision | 148 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---------------------------------------|
| Company | Sequent Medical, Inc. |
| Location | Aliso Viejo, CA, US |
| Contact | Bethany Barrett |
| 510(k) history | 4 submissions · 4 cleared · 2013-2017 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k150894/> Data sourced from FDA 510(k) public records (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. - Generated June 1, 2026