

K110980 CENTRIMAG RETURN CANNULA KITJun 20, 2011
74 days to decisionK110980 · Product code: **DWF** · Cardiovascular
Source: <https://www.510kdatabase.net/k110980/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF) |
| Date received | Apr 7, 2011 |
| Decision date | Jun 20, 2011 |
| Days to decision | 74 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Levitronix, LLC |
| Location | Waltham, MA, US |
| Contact | Lydia Sakakeeny |
| 510(k) history | 11 submissions · 11 cleared · 2003-2011 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k110980/> 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 19, 2026