

K140286 OPTease VENA CAVA FILTER AND OPTease RETRIEVAL CATHETER

Mar 7, 2014
30 days to decision

K140286 · Product code: **DTK** · Cardiovascular
Source: <https://www.510kdatabase.net/k140286/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Filter, Intravascular, Cardiovascular (DTK)
Date received	Feb 5, 2014
Decision date	Mar 7, 2014
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cordis, A Johnson & Johnson Co.
Location	Warren, NJ, US
Contact	SHAMSA KARIMI
510(k) history	4 submissions · 2 cleared · 1999-2014

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k140286/> 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 7, 2026