

**K062887 MODIFICATION TO G2 FILTER SYSTEM-FEMORAL
DELIVERY KIT, MODEL RF-310F**Oct 26, 2006
30 days to decisionK062887 · Product code: **DTK** · Cardiovascular
Source: <https://www.510kdatabase.net/k062887/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Filter, Intravascular, Cardiovascular (DTK)
Date received	Sep 26, 2006
Decision date	Oct 26, 2006
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Bard Peripheral Vascular
Location	Tempe, AZ, US
Contact	GENEVIEVE BALUTOWSKI
510(k) history	4 submissions · 4 cleared · 2003-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k062887/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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