

K113869 ORIGEN REINFORCED DUAL LUMEN CATHETERJul 26, 2012
209 days to decisionK113869 · Product code: **DWF** · Cardiovascular
Source: <https://www.510kdatabase.net/k113869/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Dec 30, 2011
Decision date	Jul 26, 2012
Days to decision	209 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Origen Biomedical, Inc.
Location	Austin, TX, US
Contact	RICHARD MARTIN
510(k) history	4 submissions · 4 cleared · 2001-2012

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