

**K070485 BOOMERANG WIRE /BOOMERANG PLUS WIRE
SYSTEM, MODEL# B PLUS WIRE 58L**Mar 22, 2007
30 days to decisionK070485 · Product code: **DXC** · Cardiovascular
Source: <https://www.510kdatabase.net/k070485/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Clamp, Vascular (DXC)
Date received	Feb 20, 2007
Decision date	Mar 22, 2007
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cardiva Medical, Inc.
Location	Pleasanton, CA, US
Contact	MICHAEL J BILLIG
510(k) history	7 submissions · 7 cleared · 2004-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k070485/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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