

K061075 BOOMERANG WIRESep 1, 2006
137 days to decisionK061075 · Product code: **DXC** · Cardiovascular
Source: <https://www.510kdatabase.net/k061075/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Clamp, Vascular (DXC)
Date received	Apr 17, 2006
Decision date	Sep 1, 2006
Days to decision	137 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k061075/> 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 9, 2026