

**K093977 ALLIGATOR-HD RETRIEVAL DEVICE (ARD-HD),  
MODEL FA-88840-XX**Apr 22, 2010  
119 days to decisionK093977 · Product code: **DQY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k093977/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Dec 24, 2009
Decision date	Apr 22, 2010
Days to decision	119 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Chestnut Medical Technologies, Inc.</b>
Location	Palo Alto, CA, US
Contact	DANIEL CHER
510(k) history	3 submissions · 3 cleared · 2005-2010

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093977/>. Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 21, 2026