FDA 510(k) Premarket Notification

K093977 ALLIGATOR-HD RETRIEVAL DEVICE (ARD-HD), MODEL FA-88840-XX

CLEARED

Apr 22, 2010
119 days to decision

K093977 · 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	**Chestnut Medical Technologies, Inc.**
Location	Palo Alto, CA, US
Contact	DANIEL CHER
510(k) history	3 submissions · 3 cleared · 2005-2010