

K151896 Houdini CatheterDec 7, 2015
150 days to decisionK151896 · Product code: **DQY** · Cardiovascular
Source: <https://www.510kdatabase.net/k151896/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jul 10, 2015
Decision date	Dec 7, 2015
Days to decision	150 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cruzar Medsystems, Inc.
Location	Braintree, MA, US
Contact	Tom Kramer
510(k) history	2 submissions · 2 cleared · 2015-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k151896/> 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 May 26, 2026