FDA 510(k) Premarket Notification

K150782 AbStats Gateway

CLEARED

Dec 14, 2015
264 days to decision

K150782 · Product code: **DQD** · Cardiovascular
Source: https://www.510kdatabase.net/k150782/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stethoscope, Electronic (DQD)
Date received	Mar 25, 2015
Decision date	Dec 14, 2015
Days to decision	264 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Gi Logic, Inc.**
Location	Pasadena, CA, US
Contact	James Beeton
510(k) history	1 submissions · 1 cleared · 2015-2015