FDA 510(k) Premarket Notification

K123551 ITCLAMP 50

CLEARED

May 14, 2013
176 days to decision

K123551 · Product code: **DXC** · Cardiovascular
Source: https://www.510kdatabase.net/k123551/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Clamp, Vascular (DXC)
Date received	Nov 19, 2012
Decision date	May 14, 2013
Days to decision	176 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Innovative Trauma Care, Inc.**
Location	San Antonio, TX, US
Contact	RICHARD WAITE
510(k) history	4 submissions · 4 cleared · 2013-2015