

**K123551 ITCLAMP 50**May 14, 2013  
176 days to decisionK123551 · Product code: **DXC** · Cardiovascular  
Source: <https://www.510kdatabase.net/k123551/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k123551/> 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 4, 2026