

**K123194 JUNCTIONAL EMERGENCY TREATMENT TOOL
MODEL 30-0088**Jan 3, 2013
84 days to decisionK123194 · Product code: **DXC** · Cardiovascular
Source: <https://www.510kdatabase.net/k123194/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Clamp, Vascular (DXC)
Date received	Oct 11, 2012
Decision date	Jan 3, 2013
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	North American Rescue Products, Inc.
Location	Greer, SC, US
Contact	WILLIAM SLEVIN
510(k) history	1 submissions · 1 cleared · 2013-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k123194/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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