FDA 510(k) Premarket Notification

K123194 JUNCTIONAL EMERGENCY TREATMENT TOOL MODEL 30-0088

CLEARED

Jan 3, 2013
84 days to decision

K123194 · 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