FDA 510(k) Premarket Notification

K073444 TRITON MEIDCAL TMED IV-SET

CLEARED

Aug 18, 2008
255 days to decision

K073444 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k073444/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Dec 7, 2007
Decision date	Aug 18, 2008
Days to decision	255 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Triton Medical, LLC**
Location	Austin, TX, US
Contact	IAN P GORDON
510(k) history	1 submissions · 1 cleared · 2008-2008