CLEARED

K122980 GLIDESHEATH SLENDER MODEL RM* ES6J10HQS, GLIDESHEATH SLENDER MODEL RM* ES6F16HQ, GLIDESHEATH SLENDER MODEL RM * RS6J10PQ

Dec 11, 2012
76 days to decision

K122980 · Product code: **DYB** · Cardiovascular
Source: https://www.510kdatabase.net/k122980/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Catheter (DYB)
Date received	Sep 26, 2012
Decision date	Dec 11, 2012
Days to decision	76 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Terumo Medical Corporation**
Location	Elkton, MD, US
Contact	DANIEL R PLONSKI
510(k) history	14 submissions · 14 cleared · 2011-2023