FDA 510(k) Premarket Notification

K150542 Terumo Pump Tubing

CLEARED

May 1, 2015
59 days to decision

K150542 · Product code: **DWE** · Cardiovascular
Source: https://www.510kdatabase.net/k150542/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Tubing, Pump, Cardiopulmonary Bypass (DWE)
Date received	Mar 3, 2015
Decision date	May 1, 2015
Days to decision	59 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Terumo Cardiovascular Systems Corporation**
Location	Elkton, MD, US
Contact	JOSHUA EWING
510(k) history	29 submissions · 29 cleared · 2002-2024