FDA 510(k) Premarket Notification

K131705 ENDO GIA RADIAL RELOAD WITH TRI-STAPLE TECHNOLOGY

CLEARED

Jul 1, 2013
20 days to decision

K131705 · Product code: **GDW** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k131705/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Staple, Implantable (GDW)
Date received	Jun 11, 2013
Decision date	Jul 1, 2013
Days to decision	20 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Covidien, Formerly US Surgical A Divison of Tyco H**
Location	North Haven, CT, US
Contact	KATHERINE ROBERTSON
510(k) history	13 submissions · 13 cleared · 2010-2013