

K131705 ENDO GIA RADIAL RELOAD WITH TRI-STAPLE TECHNOLOGY

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k131705/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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