FDA 510(k) Premarket Notification

K130717 ENDO GIA RELOAD

CLEARED

Apr 18, 2013
34 days to decision

K130717 · Product code: **GDW** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k130717/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Staple, Implantable (GDW)
Date received	Mar 15, 2013
Decision date	Apr 18, 2013
Days to decision	34 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/k130717/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 8, 2026