FDA 510(k) Premarket Notification

K130435 VERSAPORT V2 BLADELESS OPTICAL TROCAR

CLEARED

Mar 14, 2013
21 days to decision

K130435 · Product code: **GCJ** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k130435/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Laparoscope, General & Plastic Surgery (GCJ)
Date received	Feb 21, 2013
Decision date	Mar 14, 2013
Days to decision	21 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	SARAH RIZK
510(k) history	13 submissions · 13 cleared · 2010-2013

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k130435/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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