

K082659 MODIFIED ENDO STICHSep 19, 2008
7 days to decisionK082659 · Product code: **OCW** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k082659/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Endoscopic Tissue Approximation Device (OCW)
Date received	Sep 12, 2008
Decision date	Sep 19, 2008
Days to decision	7 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Covidien Lp, Formerly Registered AS United States
Location	North Haven, CT, US
Contact	ROBERT ZOTT
510(k) history	7 submissions · 7 cleared · 2008-2009

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k082659/> 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 22, 2026