

**K101797 SONICISION CORDLESS ULTRASONIC DISSECTION  
DEVICE MODEL SCD391, SCD396, SCT12, SCG, SCB, SCSK,  
SONICISION STERILIZATION T**Feb 24, 2011  
241 days to decisionK101797 · Product code: LFL · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k101797/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Ultrasonic Surgical (LFL)
Date received	Jun 28, 2010
Decision date	Feb 24, 2011
Days to decision	241 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Covidien, Formerly Valleylab, A Division of Tyco</b>
Location	Boulder, CO, US
Contact	DAVID M HORTON
510(k) history	1 submissions · 1 cleared · 2011-2011

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k101797/> 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 19, 2026