

K212301 Sonicision 7 Curved Jaw Cordless Ultrasonic Dissection DeviceNov 15, 2021
115 days to decision

K212301 · Product code: LFL · General & Plastic Surgery

Source: <https://www.510kdatabase.net/k212301/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Ultrasonic Surgical (LFL)
Date received	Jul 23, 2021
Decision date	Nov 15, 2021
Days to decision	115 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Covidien, LLC
Location	Mansfield, MA, US
Contact	Celso Duran
510(k) history	88 submissions · 85 cleared · 2010-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k212301/>. 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 May 13, 2026