

K150682 CUSA Excel+Dec 18, 2015
276 days to decisionK150682 · Product code: **LFL** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k150682/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Ultrasonic Surgical (LFL)
Date received	Mar 17, 2015
Decision date	Dec 18, 2015
Days to decision	276 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Integra Lifesciences Corp.
Location	Somerville, NJ, US
Contact	JENNIFER SIEGEL
Website	http://www.integra-ls.com/
510(k) history	29 submissions · 29 cleared · 1999-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k150682/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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