FDA 510(k) Premarket Notification

K150682 CUSA Excel+

CLEARED

Dec 18, 2015
276 days to decision

K150682 · Product code: **LFL** · General & Plastic Surgery
Source: 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