

K073322 FOX 1-980, FOX Q-1064, FOX-Q-810Feb 22, 2008
88 days to decisionK073322 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k073322/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Nov 26, 2007
Decision date	Feb 22, 2008
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	A.R.C Laser GmbH
Location	Rockville, MD, US
Contact	KIRSTEN H PAULSON
510(k) history	6 submissions · 6 cleared · 2008-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k073322/> 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 26, 2026