

K032023 W.O.M. LASER U100PLUSOct 1, 2003
93 days to decisionK032023 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k032023/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jun 30, 2003
Decision date	Oct 1, 2003
Days to decision	93 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	W.O.M. World of Medicine AG
Location	Cambridge, MA, US
Contact	SUSANNE RAAB
510(k) history	15 submissions · 15 cleared · 2002-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k032023/> 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 19, 2026