

K974641 FUTURELASE 3000/3002 ERBIUM LASER SYSTEMJun 8, 1998
175 days to decisionK974641 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k974641/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Dec 15, 1997
Decision date	Jun 8, 1998
Days to decision	175 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Buckman Co., Inc.
Location	Mchenry, IL, US
Contact	DAVID W SCHLERF
510(k) history	111 submissions · 104 cleared · 1983-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k974641/> 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 21, 2026