

K920004 PBI MULTILASE D COPPER VAPOR LASERMay 14, 1992
133 days to decisionK920004 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k920004/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Jan 2, 1992
Decision date	May 14, 1992
Days to decision	133 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Wm.I.Kern & Co., Inc.
Location	Smithtown, NY, US
Contact	WILLIAM I.KERN
510(k) history	1 submissions · 1 cleared · 1992-1992

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k920004/> 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 July 7, 2026