

K822644 RIWOLITH 2135Sep 21, 1982
20 days to decisionK822644 · Product code: **FFK** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k822644/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lithotripter, Electro-hydraulic (FFK)
Date received	Sep 1, 1982
Decision date	Sep 21, 1982
Days to decision	20 days
Third-party review	No

APPLICANT

Company	R. Wolf Medical Instruments Corp.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1981-1982

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k822644/> 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 3, 2026