FDA 510(k) Premarket Notification

K822644 RIWOLITH 2135

CLEARED

Sep 21, 1982
20 days to decision

K822644 · Product code: **FFK** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k822644/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lithotriptor, 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