

K012482 STORZ MODULITH SLK LITHOTRIPTER WITH MULTIVIEW OPTIONOct 16, 2001
75 days to decisionK012482 · Product code: **LNS** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k012482/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lithotripter, Extracorporeal Shock-wave, Urological (LNS)
Date received	Aug 2, 2001
Decision date	Oct 16, 2001
Days to decision	75 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	KARL STORZ Endoscopy-America, Inc.
Location	Mchenry, IL, US
Contact	JENNIFER PORTUGAL
510(k) history	361 submissions · 361 cleared · 1980-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k012482/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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