

**K981717 STRYKER KNIFELIGHT, STRYKER ILLUMINATED
RETRACTOR**

Jun 30, 1998
46 days to decision

K981717 · Product code: **FTD** · General & Plastic Surgery
Source: <https://www.510kdatabase.net/k981717/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lamp, Surgical (FTD)
Date received	May 15, 1998
Decision date	Jun 30, 1998
Days to decision	46 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Stryker Instruments
Location	Kalamazoo, MI, US
Contact	NICOLE PETTY
510(k) history	73 submissions · 73 cleared · 1994-2026

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k981717/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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