FDA 510(k) Premarket Notification

K981717 STRYKER KNIFELIGHT, STRYKER ILLUMINATED RETRACTOR

CLEARED

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