FDA 510(k) Premarket Notification

K896822 SURGISMOKE

CLEARED

Jan 11, 1990
38 days to decision

K896822 · Product code: **FYD** · General Hospital
Source: https://www.510kdatabase.net/k896822/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Exhaust, Surgical (FYD)
Date received	Dec 4, 1989
Decision date	Jan 11, 1990
Days to decision	38 days
Third-party review	No

APPLICANT

Company	**Spectro Industries, Inc.**
Location	Jenkintown, PA, US
Contact	DONALD HEWITT
510(k) history	2 submissions · 2 cleared · 1984-1990