FDA 510(k) Premarket Notification

K880890 THE S.A.F.E. SYSTEM

CLEARED

May 11, 1988
70 days to decision

K880890 · Product code: **FYD** · General Hospital
Source: https://www.510kdatabase.net/k880890/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Exhaust, Surgical (FYD)
Date received	Mar 2, 1988
Decision date	May 11, 1988
Days to decision	70 days
Third-party review	No

APPLICANT

Company	**Pfizer Laser Systems**
Location	Irvine, CA, US
Contact	LINDA DIBENEDETTO
510(k) history	32 submissions · 32 cleared · 1988-1993