FDA 510(k) Premarket Notification

K894429 MODIFIED CLAIMS FOR FEF(TM) END-TIDAL CO2 DETECTOR

CLEARED

Sep 18, 1989
63 days to decision

K894429 · Product code: **CCK** · Anesthesiology
Source: https://www.510kdatabase.net/k894429/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Carbon-dioxide, Gaseous-phase (CCK)
Date received	Jul 17, 1989
Decision date	Sep 18, 1989
Days to decision	63 days
Third-party review	No

APPLICANT

Company	**Fenem, Inc.**
Location	New York, NY, US
Contact	SHEILA GLENNON
510(k) history	5 submissions · 5 cleared · 1988-1989