FDA 510(k) Premarket Notification

K890436 TECHNOMED PULSOLITH LASER SYSTEM

CLEARED

Mar 23, 1989
55 days to decision

K890436 · Product code: **FFK** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k890436/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lithotriptor, Electro-hydraulic (FFK)
Date received	Jan 27, 1989
Decision date	Mar 23, 1989
Days to decision	55 days
Third-party review	No

APPLICANT

Company	**Teknomed, Inc.**
Location	Mchenry, IL, US
Contact	RICHARD HUNTER
510(k) history	7 submissions · 7 cleared · 1979-1993