FDA 510(k) Premarket Notification

K850298 ULTRALITH ULTRASONIC LITHOTRIPSY SYSTEM

CLEARED

Mar 5, 1985
40 days to decision

K850298 · Product code: **FEO** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k850298/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lithotriptor, Ultrasonic (FEO)
Date received	Jan 24, 1985
Decision date	Mar 5, 1985
Days to decision	40 days
Third-party review	No

APPLICANT

Company	**Cabot Medical Corp.**
Location	Mchenry, IL, US
Contact	DAVID E GRONOSTAJSK
510(k) history	38 submissions · 38 cleared · 1983-1996