FDA 510(k) Premarket Notification

K860571 F.M. WEIST,KG (URODYNAMIC SYSTEMS)

CLEARED

Mar 18, 1986
32 days to decision

K860571 · Product code: **FAP** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k860571/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cystometric Gas (carbon-dioxide) On Hydraulic Device (FAP)
Date received	Feb 14, 1986
Decision date	Mar 18, 1986
Days to decision	32 days
Third-party review	No

APPLICANT

Company	**F.M. Wiest USA, Inc.**
Location	Cherry Hill, NJ, US
Contact	WAYNE W DISANZA
510(k) history	12 submissions · 12 cleared · 1986-1994