FDA 510(k) Premarket Notification

K872022 UROVIEW

CLEARED

Jul 16, 1987
51 days to decision

K872022 · Product code: **KQS** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k872022/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Cystometric, Non-electric And Accessories (KQS)
Date received	May 26, 1987
Decision date	Jul 16, 1987
Days to decision	51 days
Third-party review	No

APPLICANT

Company	**Oec-Diasonics, Inc.**
Location	Salt Lake City, UT, US
Contact	JOHN W TOLHURST
510(k) history	8 submissions · 8 cleared · 1985-1993