

K872022 UROVIEWJul 16, 1987
51 days to decisionK872022 · Product code: **KQS** · Gastroenterology & UrologySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k872022/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 20, 2026