

K970982 T-RX RETRIEVAL FORCEPSMay 9, 1997
52 days to decisionK970982 · Product code: **OCZ** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k970982/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscopic Grasping/cutting Instrument, Non-powered (OCZ)
Date received	Mar 18, 1997
Decision date	May 9, 1997
Days to decision	52 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Tri-Med Specialties, Inc.
Location	Overland Park, KS, US
Contact	MIKE KNOTH
510(k) history	14 submissions · 12 cleared · 1985-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k970982/> 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 July 1, 2026