

K926040 HELI-TECH RETRIEVAL BASKETDec 23, 1992
22 days to decisionK926040 · Product code: **LQR** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k926040/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dislodger, Stone, Biliary (LQR)
Date received	Dec 1, 1992
Decision date	Dec 23, 1992
Days to decision	22 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cape Tech Medical Products
Location	Falmouth, MA, US
Contact	AL BICCUM
510(k) history	3 submissions · 3 cleared · 1992-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k926040/> 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 2, 2026