

K960004 CELL RECOVERY SYSTEMMar 20, 1996
78 days to decisionK960004 · Product code: **FDX** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k960004/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Endoscopic Cytology Brush (FDX)
Date received	Jan 2, 1996
Decision date	Mar 20, 1996
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medical Device Technologies, Inc.
Location	Gainesville, FL, US
Contact	M. LEE HULSEBUS
510(k) history	46 submissions · 46 cleared · 1992-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k960004/> 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