FDA 510(k) Premarket Notification

K960004 CELL RECOVERY SYSTEM

CLEARED

Mar 20, 1996
78 days to decision

K960004 · 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	Gainsville, FL, US
Contact	M. LEE HULSEBUS
510(k) history	46 submissions · 46 cleared · 1992-2010