FDA 510(k) Premarket Notification

K893390 MODIFIED PRISTINE CULTURE CATHETER

CLEARED

Sep 7, 1989
129 days to decision

K893390 · Product code: **KOD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k893390/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Urological (KOD)
Date received	May 1, 1989
Decision date	Sep 7, 1989
Days to decision	129 days
Third-party review	No

APPLICANT

Company	**Hml Medical, Inc.**
Location	Irvine, CA, US
Contact	ERIC M LOVGREN
510(k) history	1 submissions · 1 cleared · 1989-1989