FDA 510(k) Premarket Notification

K875171 J-CATH(TM)

CLEARED

Apr 6, 1988
111 days to decision

K875171 · Product code: **KNT** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k875171/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Gastrointestinal (and Accessories) (KNT)
Date received	Dec 17, 1987
Decision date	Apr 6, 1988
Days to decision	111 days
Third-party review	No

APPLICANT

Company	**Hdc Corp.**
Location	Walker, MI, US
Contact	RONALD L COLEMAN
510(k) history	30 submissions · 29 cleared · 1983-2007