FDA 510(k) Premarket Notification

K800505 URETHRAL CATHETER

CLEARED

Apr 8, 1980
35 days to decision

K800505 · Product code: **KOD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k800505/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Urological (KOD)
Date received	Mar 4, 1980
Decision date	Apr 8, 1980
Days to decision	35 days
Third-party review	No

APPLICANT

Company	**Medi-Craft , Ltd.**
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1980-1986