FDA 510(k) Premarket Notification

K874293 ACKRAD V/CUG VOIDING CYSTOURETHROGRAPHY SET

CLEARED

Dec 29, 1987
70 days to decision

K874293 · Product code: **KOD** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k874293/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Urological (KOD)
Date received	Oct 20, 1987
Decision date	Dec 29, 1987
Days to decision	70 days
Third-party review	No

APPLICANT

Company	**Ackrad Laboratories**
Location	Mchenry, IL, US
Contact	BERNARD ACKERMAN
510(k) history	42 submissions · 41 cleared · 1979-2002