

**K874293 ACKRAD V/CUG VOIDING CYSTOURETHROGRAPHY SET**Dec 29, 1987  
70 days to decisionK874293 · 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	<b>Ackrad Laboratories</b>
Location	Mchenry, IL, US
Contact	BERNARD ACKERMAN
510(k) history	42 submissions · 41 cleared · 1979-2002

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k874293/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026