

**K881680 ACKRAD H/S PROCEDURE TRAY**Jul 15, 1988  
87 days to decisionK881680 · Product code: **HIG** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k881680/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SD
Submission type	Traditional
Device classification	Insufflator, Hysteroscopic (HIG)
Date received	Apr 19, 1988
Decision date	Jul 15, 1988
Days to decision	87 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881680/>, Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 26, 2026