

**K884696 RESUBMITTED ARTIFICIAL INSEMINATION  
INSTRUMENT SET**Jan 27, 1989  
79 days to decisionK884696 · Product code: **HDR** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k884696/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cap, Cervical (HDR)
Date received	Nov 9, 1988
Decision date	Jan 27, 1989
Days to decision	79 days
Third-party review	No

**APPLICANT**

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Company	<b>Laboratoire Ccd C/O Washington Regulatory Services</b>
Location	Ringoes, NJ, US
Contact	RANDOLPH L COOKE
510(k) history	4 submissions · 4 cleared · 1988-1989

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k884696/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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