FDA 510(k) Premarket Notification

K884696 RESUBMITTED ARTIFICIAL INSEMINATION INSTRUMENT SET

CLEARED

Jan 27, 1989
79 days to decision

K884696 · Product code: **HDR** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k884696/

SUBMISSION DETAILS

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

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