

K831669 AXCAN SPERM CUPJul 12, 1983
50 days to decisionK831669 · Product code: **HDR** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k831669/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cap, Cervical (HDR)
Date received	May 23, 1983
Decision date	Jul 12, 1983
Days to decision	50 days
Third-party review	No

APPLICANT

Company	Axcan Scientific Corp.
Location	Walker, MI, US
510(k) history	5 submissions · 5 cleared · 1982-1984

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Device record: <https://www.510kdatabase.net/k831669/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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