

**K831212 ENDOSCANN**Jun 15, 1984  
428 days to decisionK831212 · Product code: **HFF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k831212/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Aspirator, Endometrial (HFF)
Date received	Apr 14, 1983
Decision date	Jun 15, 1984
Days to decision	428 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k831212/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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