

**K973850 ALEXANDER MANUFACTURING CO.  
RECHARGEABLE BATTERY #M007**Nov 12, 1997  
35 days to decisionK973850 · Product code: **HFO** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k973850/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Pressure, Intrauterine (HFO)
Date received	Oct 8, 1997
Decision date	Nov 12, 1997
Days to decision	35 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Alexander Mfg. Co.</b>
Location	Mason City, IA, US
Contact	KEN HEIMENDINGER
510(k) history	39 submissions · 37 cleared · 1997-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k973850/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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