

**K823937 SEMINAL COLLECTION DEVICE**Feb 24, 1983  
57 days to decisionK823937 · Product code: **HIS** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k823937/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Condom (HIS)
Date received	Dec 29, 1982
Decision date	Feb 24, 1983
Days to decision	57 days
Third-party review	No

**APPLICANT**

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Company	<b>Health Development Corp.</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1980-1983

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

Device record: <https://www.510kdatabase.net/k823937/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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