

**K810832 FORECASTER**Jun 24, 1981  
91 days to decisionK810832 · Product code: **LHD** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k810832/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Fertility Diagnostic, Proceptive (LHD)
Date received	Mar 25, 1981
Decision date	Jun 24, 1981
Days to decision	91 days
Third-party review	No

**APPLICANT**

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Company	<b>Forecaster Co.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k810832/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 30, 2026