

**K905115 BIOVISION ENDOCULAR PROBE**Feb 11, 1991  
90 days to decisionK905115 · Product code: **HGF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k905115/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Source, Abortion Unit, Vacuum (HGF)
Date received	Nov 13, 1990
Decision date	Feb 11, 1991
Days to decision	90 days
Third-party review	No

**APPLICANT**

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Company	<b>Douglas James Donaldson Compliance Services</b>
Location	Oakland, CA, US
Contact	DOUGLAS DONALDSON
510(k) history	5 submissions · 5 cleared · 1990-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k905115/>; 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