

**K891465 OBSTETRIC CUP (MODIFICATION)**May 11, 1989  
57 days to decisionK891465 · Product code: **HDB** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k891465/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Extractor, Vacuum, Fetal (HDB)
Date received	Mar 15, 1989
Decision date	May 11, 1989
Days to decision	57 days
Third-party review	No

**APPLICANT**

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Company	<b>Go Medical Industries Pty. , Ltd.</b>
Location	Walker, MI, US
Contact	GEORGE O'NEIL
510(k) history	11 submissions · 11 cleared · 1984-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k891465/>; 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