

K840384 DIAGNOSTIC DILATATION & MENSTRUAL KITApr 30, 1984
91 days to decisionK840384 · Product code: **HHI** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k840384/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Abortion, Vacuum (HHI)
Date received	Jan 30, 1984
Decision date	Apr 30, 1984
Days to decision	91 days
Third-party review	No

APPLICANT

Company	Intl. Medical Technologies, Inc.
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1984-1984

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Device record: <https://www.510kdatabase.net/k840384/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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