

**K840766 JANACEK TUBO-UTERINE IMPLANTATION SET**Apr 23, 1984  
60 days to decisionK840766 · Product code: **HTD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k840766/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Forceps (HTD)
Date received	Feb 23, 1984
Decision date	Apr 23, 1984
Days to decision	60 days
Third-party review	No

**APPLICANT**

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Company	<b>Cimed Intl., Inc.</b>
Location	Mchenry, IL, US
Contact	JAMES L KENWORTHY
510(k) history	3 submissions · 3 cleared · 1984-1984

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k840766/>; 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 July 8, 2026