

**K905240 VESSELOOPS**Feb 6, 1991  
78 days to decisionK905240 · Product code: **KDC** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k905240/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Surgical, Disposable (KDC)
Date received	Nov 20, 1990
Decision date	Feb 6, 1991
Days to decision	78 days
Third-party review	No

**APPLICANT**

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Company	<b>Sparta Surgical Corp.</b>
Location	Hayward, CA, US
Contact	CARLETON F KIMBER
510(k) history	7 submissions · 6 cleared · 1989-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k905240/>, 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