

**K901718 KLEIN LAMPREY-CANNULA(TM)**May 24, 1990  
41 days to decisionK901718 · Product code: **GEA** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k901718/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Surgical, General & Plastic Surgery (GEA)
Date received	Apr 13, 1990
Decision date	May 24, 1990
Days to decision	41 days
Third-party review	No

**APPLICANT**

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Company	<b>Jeff Klein Surgical, Inc.</b>
Location	San Clemente, CA, US
Contact	JACQUELINE KLEIN
510(k) history	5 submissions · 5 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k901718/>; 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 7, 2026