

**K854890 HOOK PROBE**May 2, 1986  
147 days to decisionK854890 · Product code: **KDC** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k854890/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Surgical, Disposable (KDC)
Date received	Dec 6, 1985
Decision date	May 2, 1986
Days to decision	147 days
Third-party review	No

**APPLICANT**

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Company	<b>Euro-Med Intl.</b>
Location	Danville, CA, US
Contact	DONALD HOLSTEN
510(k) history	16 submissions · 16 cleared · 1985-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k854890/>, 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 May 22, 2026