

**K890234 SUTURE CLIP REMOVER**Jan 27, 1989  
9 days to decisionK890234 · Product code: **HTD** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k890234/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Forceps (HTD)
Date received	Jan 18, 1989
Decision date	Jan 27, 1989
Days to decision	9 days
Third-party review	No

**APPLICANT**

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Company	<b>Kinetic Medical Products</b>
Location	Erie, PA, US
Contact	JAMES I LAUGHNER
510(k) history	63 submissions · 63 cleared · 1989-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890234/>; 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