

**K922878 MODEL GT1ST CRYOSURGICAL INSTRUMENT**Sep 2, 1992  
79 days to decisionK922878 · Product code: **GEH** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k922878/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Cryosurgical, Accessories (GEH)
Date received	Jun 15, 1992
Decision date	Sep 2, 1992
Days to decision	79 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Gyne-Tech Instrument Corp.</b>
Location	Burbank, CA, US
Contact	KERMIT FLOYD
510(k) history	3 submissions · 3 cleared · 1992-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k922878/> 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