

K920040 INTRADUCERFeb 28, 1992
53 days to decisionK920040 · Product code: **GCJ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k920040/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Laparoscope, General & Plastic Surgery (GCJ)
Date received	Jan 6, 1992
Decision date	Feb 28, 1992
Days to decision	53 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Taut, Inc.
Location	Walker, MI, US
Contact	KENSETH
510(k) history	16 submissions · 16 cleared · 1983-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k920040/> Data sourced from FDA 510(k) public records (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. - Generated June 29, 2026