

**K924981 OPERATIVE CHOLANGIOGRAM CATHETER**Apr 6, 1993  
187 days to decisionK924981 · Product code: **GBZ** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k924981/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cholangiography (GBZ)
Date received	Oct 1, 1992
Decision date	Apr 6, 1993
Days to decision	187 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k924981/> 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 29, 2026