

K001807 TISSUE MARKERMay 3, 2001
322 days to decisionK001807 · Product code: **GDW** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k001807/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Staple, Implantable (GDW)
Date received	Jun 15, 2000
Decision date	May 3, 2001
Days to decision	322 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Advanced Uroscience, Inc.
Location	St, Paul, MN, US
Contact	KAREN E PETERSON
510(k) history	5 submissions · 5 cleared · 1998-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k001807/> 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 30, 2026