

K060268 PEDIATRIC URINARY CATHETERFeb 24, 2006
23 days to decisionK060268 · Product code: **GBM** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k060268/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Urethral (GBM)
Date received	Feb 1, 2006
Decision date	Feb 24, 2006
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Clay Kennard
Location	Oklahoma City, OK, US
Contact	CLAY KENNARD
510(k) history	1 submissions · 1 cleared · 2006-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k060268/> 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