

K063233 DELTACUT (APPARATUS AND CANNULA)Apr 30, 2007
187 days to decisionK063233 · Product code: **KNW** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k063233/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Oct 25, 2006
Decision date	Apr 30, 2007
Days to decision	187 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	PAJUNK GmbH Medizintechnologie
Location	Geisingen, DE
Contact	CHRISTIAN QUASS
510(k) history	44 submissions · 44 cleared · 2006-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k063233/> 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 28, 2026