

K061021 ENTAKE STANDARD AND SAFETY PEG SYSTEMApr 28, 2006
15 days to decisionK061021 · Product code: **KNT** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k061021/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Gastrointestinal (and Accessories) (KNT)
Date received	Apr 13, 2006
Decision date	Apr 28, 2006
Days to decision	15 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Conmedcorp
Location	Dayton, OH, US
Contact	BETH A ZIS
510(k) history	92 submissions · 92 cleared · 1981-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k061021/> 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 May 27, 2026