FDA 510(k) Premarket Notification

K061021 ENTAKE STANDARD AND SAFETY PEG SYSTEM

CLEARED

Apr 28, 2006
15 days to decision

K061021 · Product code: **KNT** · Gastroenterology & Urology
Source: 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