FDA 510(k) Premarket Notification

K133599 CORGRIP

CLEARED

Mar 26, 2014
121 days to decision

K133599 · Product code: **KNT** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k133599/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubes, Gastrointestinal (and Accessories) (KNT)
Date received	Nov 25, 2013
Decision date	Mar 26, 2014
Days to decision	121 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Corpak Medsystems**
Location	Buffalo Grove, IL, US
Contact	STEPHANIE WASIELEWSKI
510(k) history	15 submissions · 15 cleared · 1982-2014