FDA 510(k) Premarket Notification

K142792 AbbVie NJ

CLEARED

Jan 14, 2015
110 days to decision

K142792 · Product code: **KNT** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k142792/

SUBMISSION DETAILS

Decision	Substantially Equivalent - K
Submission type	Traditional
Device classification	Tubes, Gastrointestinal (and Accessories) (KNT)
Date received	Sep 26, 2014
Decision date	Jan 14, 2015
Days to decision	110 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**AbbVie, Inc.**
Location	North Chicago, IL, US
Contact	Katherine Wortley
510(k) history	6 submissions · 4 cleared · 2014-2015