

K142792 AbbVie NJJan 14, 2015
110 days to decisionK142792 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k142792/> 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 5, 2026