FDA 510(k) Premarket Notification

K132314 MEDLINE ANTI-REFLUX VALVE

CLEARED

Sep 18, 2013
55 days to decision

K132314 · Product code: **FEG** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k132314/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Double Lumen For Intestinal Decompression And/or Intubation (FEG)
Date received	Jul 25, 2013
Decision date	Sep 18, 2013
Days to decision	55 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Medline Industries, Inc.**
Location	Mchenry, IL, US
Contact	MATT CLAUSEN
510(k) history	238 submissions · 234 cleared · 1977-2024