

K132314 MEDLINE ANTI-REFLUX VALVESep 18, 2013
55 days to decisionK132314 · Product code: **FEG** · Gastroenterology & UrologySource: <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

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