FDA 510(k) Premarket Notification

K880850 KEITH NASOGASTRIC SUMP TUBE W/ANTI-REFLUX VALVE

CLEARED

May 26, 1988
86 days to decision

K880850 · Product code: **FEG** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k880850/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Double Lumen For Intestinal Decompression And/or Intubation (FEG)
Date received	Mar 1, 1988
Decision date	May 26, 1988
Days to decision	86 days
Third-party review	No

APPLICANT

Company	**Kmb Medical Products Co.**
Location	Hazelwood, MO, US
Contact	C. W KEITH
510(k) history	1 submissions · 1 cleared · 1988-1988