

K880850 KEITH NASOGASTRIC SUMP TUBE W/ANTI-REFLUX VALVEMay 26, 1988
86 days to decisionK880850 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k880850/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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