

K812305 MONTG. ESOPHAGEAL -SALIVARY BYPASS- TUBESep 8, 1981
22 days to decisionK812305 · Product code: **KCF** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k812305/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Esophageal, Ent (KCF)
Date received	Aug 17, 1981
Decision date	Sep 8, 1981
Days to decision	22 days
Third-party review	No

APPLICANT

Company	Boston Medical Products, Inc.
Location	Mchenry, IL, US
510(k) history	33 submissions · 33 cleared · 1980-2009

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Device record: <https://www.510kdatabase.net/k812305/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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