

**K962688 MONTGOMERY RADIOPAQUE SALIVARY BYPASS
TUBE**Jul 25, 1996
14 days to decisionK962688 · Product code: **KCF** · Ear, Nose, Throat
Source: <https://www.510kdatabase.net/k962688/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Esophageal, Ent (KCF)
Date received	Jul 11, 1996
Decision date	Jul 25, 1996
Days to decision	14 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k962688/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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