FDA 510(k) Premarket Notification

K962688 MONTGOMERY RADIOPAQUE SALIVARY BYPASS TUBE

CLEARED

Jul 25, 1996
14 days to decision

K962688 · 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