FDA 510(k) Premarket Notification

K812305 MONTG. ESOPHAGEAL -SALIVARY BYPASS- TUBE

CLEARED

Sep 8, 1981
22 days to decision

K812305 · Product code: **KCF** · Ear, Nose, Throat
Source: 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