FDA 510(k) Premarket Notification

K800623 JACKSON ESOPHAGEAL BOUGIE, X-RAY OPAQUE

CLEARED

Apr 24, 1980
36 days to decision

K800623 · Product code: **KCD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k800623/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bougie, Esophageal, Ent (KCD)
Date received	Mar 19, 1980
Decision date	Apr 24, 1980
Days to decision	36 days
Third-party review	No

APPLICANT

Company	**Pilling Co.**
Location	Mchenry, IL, US
Website	https://www.pilling.com
510(k) history	39 submissions · 39 cleared · 1976-1994

Pilling Co. is a medical device manufacturer based in McHenry, US, specializing in surgical instrumentation and specialized medical devices. The company has received FDA 510(k) clearances from total submissions, with clearances spanning from 1976 to 1994. Pilling Co. focused primarily on General & Plastic Surgery devices, alongside offerings in Cardiovascular, Dental, Gastroenterology & Urology, and Anesthesiology categories. Notable cleared devices include laparoscopic instrumentation, microlaryngeal forceps, vascular instruments, and coronary perfusion catheters. This c...