FDA 510(k) Premarket Notification

K812289 CELESTIN DILATOR

CLEARED

Sep 1, 1981
19 days to decision

K812289 · Product code: **KNQ** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k812289/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Esophageal (KNQ)
Date received	Aug 13, 1981
Decision date	Sep 1, 1981
Days to decision	19 days
Third-party review	No

APPLICANT

Company	**Inmed Corp.**
Location	Mchenry, IL, US
510(k) history	32 submissions · 32 cleared · 1981-1986