FDA 510(k) Premarket Notification

K822260 TRACHEAL DILATORS

CLEARED

Sep 2, 1982
35 days to decision

K822260 · Product code: **KCG** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k822260/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Tracheal (KCG)
Date received	Jul 29, 1982
Decision date	Sep 2, 1982
Days to decision	35 days
Third-party review	No

APPLICANT

Company	**Kelleher Corp.**
Location	Mchenry, IL, US
510(k) history	94 submissions · 94 cleared · 1982-1983