

K822260 TRACHEAL DILATORSSep 2, 1982
35 days to decisionK822260 · Product code: **KCG** · Ear, Nose, ThroatSource: <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

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Device record: <https://www.510kdatabase.net/k822260/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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