FDA 510(k) Premarket Notification

K822201 EAR CANNULAS

CLEARED

Oct 6, 1982
75 days to decision

K822201 · Product code: **FGY** · General & Plastic Surgery
Source: https://www.510kdatabase.net/k822201/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Injection (FGY)
Date received	Jul 23, 1982
Decision date	Oct 6, 1982
Days to decision	75 days
Third-party review	No

APPLICANT

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