

**K822201 EAR CANNULAS**Oct 6, 1982  
75 days to decisionK822201 · Product code: **FGY** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k822201/>**SUBMISSION DETAILS**

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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**

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

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Device record: <https://www.510kdatabase.net/k822201/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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