

K781631 ECHO-OILOct 10, 1978
15 days to decisionK781631 · Product code: **EMX** · Radiology
Source: <https://www.510kdatabase.net/k781631/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Balloon, Epistaxis (EMX)
Date received	Sep 25, 1978
Decision date	Oct 10, 1978
Days to decision	15 days
Third-party review	No

APPLICANT

Company	Echo Laboratories, Inc.
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1978-1982

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Device record: <https://www.510kdatabase.net/k781631/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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