FDA 510(k) Premarket Notification

K781631 ECHO-OIL

CLEARED

Oct 10, 1978
15 days to decision

K781631 · 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