

**K800175 NASOSTAT**Feb 21, 1980  
24 days to decisionK800175 · Product code: **EMX** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k800175/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Balloon, Epistaxis (EMX)
Date received	Jan 28, 1980
Decision date	Feb 21, 1980
Days to decision	24 days
Third-party review	No

**APPLICANT**

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Company	<b>Edward Weck, Inc.</b>
Location	Mchenry, IL, US
510(k) history	140 submissions · 140 cleared · 1976-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k800175/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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