

**K791680 POPE UMBRELLA TUBE**Nov 13, 1979  
78 days to decisionK791680 · Product code: **ETD** · Ear, Nose, ThroatSource: <https://www.510kdatabase.net/k791680/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tube, Tympanostomy (ETD)
Date received	Aug 27, 1979
Decision date	Nov 13, 1979
Days to decision	78 days
Third-party review	No

**APPLICANT**

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Company	<b>Xomed, Inc.</b>
Location	Arnold, MO, US
510(k) history	82 submissions · 81 cleared · 1976-2000

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

Device record: <https://www.510kdatabase.net/k791680/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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