FDA 510(k) Premarket Notification

K791680 POPE UMBRELLA TUBE

CLEARED

Nov 13, 1979
78 days to decision

K791680 · Product code: **ETD** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k791680/

SUBMISSION DETAILS

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

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