FDA 510(k) Premarket Notification

K792018 SILICONE RUBBER-BI NASOPHARYNGEAL AIR

CLEARED

Nov 13, 1979
35 days to decision

K792018 · Product code: **BTQ** · Anesthesiology
Source: https://www.510kdatabase.net/k792018/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Airway, Nasopharyngeal (BTQ)
Date received	Oct 9, 1979
Decision date	Nov 13, 1979
Days to decision	35 days
Third-party review	No

APPLICANT

Company	**Vesta, Inc.**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1979-1983