

**K792018 SILICONE RUBBER-BI NASOPHARYNGEAL AIR**Nov 13, 1979  
35 days to decisionK792018 · Product code: **BTQ** · AnesthesiologySource: <https://www.510kdatabase.net/k792018/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k792018/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 27, 2026