

**K882219 P.E.E.P. VALVE**Aug 30, 1988  
96 days to decisionK882219 · Product code: **BYE** · AnesthesiologySource: <https://www.510kdatabase.net/k882219/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Attachment, Breathing, Positive End Expiratory Pressure (BYE)
Date received	May 26, 1988
Decision date	Aug 30, 1988
Days to decision	96 days
Third-party review	No

**APPLICANT**

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Company	<b>Life Support Products, Inc.</b>
Location	Anaheim, CA, US
Contact	TOM FANGROW
510(k) history	20 submissions · 20 cleared · 1985-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882219/>; 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 26, 2026