

**K932799 SINGLE USE MANUAL RESUSCITATOR PEEP VALVE**Sep 3, 1993  
86 days to decisionK932799 · Product code: **BYE** · Anesthesiology  
Source: <https://www.510kdatabase.net/k932799/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Attachment, Breathing, Positive End Expiratory Pressure (BYE)
Date received	Jun 9, 1993
Decision date	Sep 3, 1993
Days to decision	86 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Engineered Medical Systems</b>
Location	Indianapolis, IN, US
Contact	JEFF QUINN
510(k) history	22 submissions · 22 cleared · 1987-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k932799/> 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 25, 2026