

**K922537 DURALIFE**Dec 4, 1992  
190 days to decisionK922537 · Product code: **BZA** · Anesthesiology  
Source: <https://www.510kdatabase.net/k922537/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Connector, Airway (extension) (BZA)
Date received	May 28, 1992
Decision date	Dec 4, 1992
Days to decision	190 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Dhd Diemolding Healthcare Div.</b>
Location	Canastota, NY, US
Contact	JEAN WALLACE
510(k) history	11 submissions · 11 cleared · 1991-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k922537/> 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