

**K902141 ANOLIFT - PORTABLE & ANOLIFT - GANTRY**Sep 12, 1990  
120 days to decisionK902141 · Product code: **FSA** · General HospitalSource: <https://www.510kdatabase.net/k902141/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Lift, Patient, Non-ac-powered (FSA)
Date received	May 15, 1990
Decision date	Sep 12, 1990
Days to decision	120 days
Third-party review	No

**APPLICANT**

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Company	<b>Anodyne Corp.</b>
Location	Brighton, MI, US
Contact	DONALD D MERRY
510(k) history	1 submissions · 1 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902141/>, 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 July 7, 2026