

**K820248 RODDER VOLUME VENTILATOR**Apr 16, 1982  
77 days to decisionK820248 · Product code: **CBK** · AnesthesiologySource: <https://www.510kdatabase.net/k820248/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Continuous, Facility Use (CBK)
Date received	Jan 29, 1982
Decision date	Apr 16, 1982
Days to decision	77 days
Third-party review	No

**APPLICANT**

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Company	<b>Pharmaquest Corp.</b>
Location	Walker, MI, US
510(k) history	14 submissions · 14 cleared · 1981-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k820248/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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