

**K014187 AUTOVAC ORTHOPEDIC AUTOTRANSFUSION SYSTEM**Feb 6, 2002  
47 days to decisionK014187 · Product code: **CAC** · Anesthesiology  
Source: <https://www.510kdatabase.net/k014187/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Apparatus, Autotransfusion (CAC)
Date received	Dec 21, 2001
Decision date	Feb 6, 2002
Days to decision	47 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Boehringer Laboratories</b>
Location	Mchenry, IL, US
Contact	JOHN KARPOWICZ
Website	<a href="http://www.boehringerlabs.com">http://www.boehringerlabs.com</a>
510(k) history	38 submissions · 38 cleared · 1976-2024

Boehringer Laboratories is a family-owned American medical technology company headquartered in Phoenixville, Pennsylvania, with operations in McHenry, US. The company specializes in respiratory therapy and minimally invasive surgical devices. Boehringer Laboratories has maintained a strong FDA 510(k) regulatory record since 1976. The company has received FDA 510(k) clearances from total submissions, with no denied submissions. Recent clearances span 2024, demonstrating continued active development. The company's portfolio focuses primarily on anesthesiology devices, inclu...

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

Device record: <https://www.510kdatabase.net/k014187/>. Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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