FDA 510(k) Premarket Notification

K853155 BOEHRINGER LAB SUCTION INTERRUPTER

CLEARED

Oct 16, 1985
79 days to decision

K853155 · Product code: **KDP** · General Hospital
Source: https://www.510kdatabase.net/k853155/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Regulator, Vacuum (KDP)
Date received	Jul 29, 1985
Decision date	Oct 16, 1985
Days to decision	79 days
Third-party review	No

APPLICANT

Company	**Boehringer Laboratories**
Location	Mchenry, IL, US
Contact	JOHN BOEHRINGER
Website	http://www.boehringerlabs.com
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...