FDA 510(k) Premarket Notification

K832868 RESUSCITATION VALVE

CLEARED

Sep 26, 1983
33 days to decision

K832868 · Product code: **CBP** · Anesthesiology
Source: https://www.510kdatabase.net/k832868/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Valve, Non-rebreathing (CBP)
Date received	Aug 24, 1983
Decision date	Sep 26, 1983
Days to decision	33 days
Third-party review	No

APPLICANT

Company	**Boehringer Laboratories**
Location	Mchenry, IL, US
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...