

**K983631 AUTO-PEEP MEASUREMENT DEVICE, MODEL # KC
9-P**Jul 1, 1999
259 days to decisionK983631 · Product code: **CBP** · Anesthesiology
Source: <https://www.510kdatabase.net/k983631/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Valve, Non-rebreathing (CBP)
Date received	Oct 15, 1998
Decision date	Jul 1, 1999
Days to decision	259 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Instrumentation Industries, Inc.
Location	Mchenry, IL, US
Contact	EDWARD C HOREY, JR.
510(k) history	45 submissions · 44 cleared · 1977-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k983631/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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