FDA 510(k) Premarket Notification

K983631 AUTO-PEEP MEASUREMENT DEVICE, MODEL # KC 9-P

CLEARED

Jul 1, 1999
259 days to decision

K983631 · 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