

K030943 MODIFICATION TO HIOXApr 22, 2003
27 days to decisionK030943 · Product code: **CBP** · Anesthesiology
Source: <https://www.510kdatabase.net/k030943/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Valve, Non-rebreathing (CBP)
Date received	Mar 26, 2003
Decision date	Apr 22, 2003
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Sensor Medics Corp.
Location	Walker, MI, US
Contact	EARL W DRAPER
510(k) history	26 submissions · 26 cleared · 1984-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k030943/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 25, 2026