FDA 510(k) Premarket Notification

K030943 MODIFICATION TO HIOX

CLEARED

Apr 22, 2003
27 days to decision

K030943 · 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