FDA 510(k) Premarket Notification

K960587 CPROTECTOR 2000

CLEARED

Jul 21, 1997
525 days to decision

K960587 · Product code: **CBP** · Anesthesiology
Source: https://www.510kdatabase.net/k960587/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Valve, Non-rebreathing (CBP)
Date received	Feb 12, 1996
Decision date	Jul 21, 1997
Days to decision	525 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Certified Safety Mfg., Inc.**
Location	Kansas City, MO, US
Contact	HOWARD GERSON
510(k) history	17 submissions · 14 cleared · 1994-1997