FDA 510(k) Premarket Notification

K953414 CPR MICROMASK WITH OXYGEN INLET

CLEARED

Mar 20, 1996
244 days to decision

K953414 · Product code: **CBP** · Anesthesiology
Source: https://www.510kdatabase.net/k953414/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Valve, Non-rebreathing (CBP)
Date received	Jul 20, 1995
Decision date	Mar 20, 1996
Days to decision	244 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Plasco, Inc.**
Location	Mchenry, IL, US
Contact	GARY S BOTSFORD
510(k) history	16 submissions · 16 cleared · 1983-1998

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k953414/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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