FDA 510(k) Premarket Notification

K943402 K.P.R. (KING PERSONAL RESUSCITATOR)

CLEARED

Aug 19, 1994
36 days to decision

K943402 · Product code: **CBP** · Anesthesiology
Source: https://www.510kdatabase.net/k943402/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Valve, Non-rebreathing (CBP)
Date received	Jul 14, 1994
Decision date	Aug 19, 1994
Days to decision	36 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**King Systems Corp.**
Location	Noblesville, IN, US
Contact	MICHAEL A LEAGRE
510(k) history	18 submissions · 18 cleared · 1988-2004

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k943402/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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