FDA 510(k) Premarket Notification

K791850 ONE-WAY VALVE-FEMALE, M-503

CLEARED

Oct 1, 1979
12 days to decision

K791850 · Product code: **CBP** · Anesthesiology
Source: https://www.510kdatabase.net/k791850/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Valve, Non-rebreathing (CBP)
Date received	Sep 19, 1979
Decision date	Oct 1, 1979
Days to decision	12 days
Third-party review	No

APPLICANT

Company	**Mem Medical, Inc.**
Location	Mchenry, IL, US
510(k) history	27 submissions · 27 cleared · 1979-1979