FDA 510(k) Premarket Notification

K802231 AIR VIVA II

CLEARED

Oct 10, 1980
25 days to decision

K802231 · Product code: **BTM** · Anesthesiology
Source: https://www.510kdatabase.net/k802231/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Emergency, Manual (resuscitator) (BTM)
Date received	Sep 15, 1980
Decision date	Oct 10, 1980
Days to decision	25 days
Third-party review	No

APPLICANT

Company	**General Medical Co.**
Location	Mchenry, IL, US
510(k) history	15 submissions · 14 cleared · 1977-1993