FDA 510(k) Premarket Notification

K780558 BOTTLE BREATHING DEVICE

CLEARED

May 26, 1978
51 days to decision

K780558 · Product code: **BYO** · Anesthesiology
Source: https://www.510kdatabase.net/k780558/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bottle, Blow (BYO)
Date received	Apr 5, 1978
Decision date	May 26, 1978
Days to decision	51 days
Third-party review	No

APPLICANT

Company	**B & F Medical Products, Inc.**
Location	Mchenry, IL, US
510(k) history	18 submissions · 18 cleared · 1976-1994