

K780558 BOTTLE BREATHING DEVICEMay 26, 1978
51 days to decisionK780558 · Product code: **BYO** · AnesthesiologySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k780558/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026