

K770619 EXERCISOR, RESPIRATORYApr 7, 1977
6 days to decisionK770619 · Product code: **BWF** · AnesthesiologySource: <https://www.510kdatabase.net/k770619/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Therapeutic (incentive) (BWF)
Date received	Apr 1, 1977
Decision date	Apr 7, 1977
Days to decision	6 days
Third-party review	No

APPLICANT

Company	Cutter Laboratories, Inc.
Location	Mchenry, IL, US
Website	https://www.bayer.com
510(k) history	39 submissions · 39 cleared · 1976-1986

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k770619/>; 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 22, 2026