

K771175 RESUSCITATOR, MS30Aug 2, 1977
34 days to decisionK771175 · Product code: **BTM** · AnesthesiologySource: <https://www.510kdatabase.net/k771175/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Emergency, Manual (resuscitator) (BTM)
Date received	Jun 29, 1977
Decision date	Aug 2, 1977
Days to decision	34 days
Third-party review	No

APPLICANT

Company	Mckesson Corp.
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1976-1977

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k771175/> 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 28, 2026