

K840349 MARK III RESUSCITATORFeb 10, 1984
15 days to decisionK840349 · Product code: **BTM** · AnesthesiologySource: <https://www.510kdatabase.net/k840349/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Emergency, Manual (resuscitator) (BTM)
Date received	Jan 26, 1984
Decision date	Feb 10, 1984
Days to decision	15 days
Third-party review	No

APPLICANT

Company	Ambu, Inc.
Location	Walker, MI, US
510(k) history	33 submissions · 33 cleared · 1984-2005

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k840349/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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