

K802231 AIR VIVA IIOct 10, 1980
25 days to decisionK802231 · Product code: **BTM** · AnesthesiologySource: <https://www.510kdatabase.net/k802231/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Emergency, Manual (resuscitator) (BTM)
Date received	Sep 15, 1980
Decision date	Oct 10, 1980
Days to decision	25 days
Third-party review	No

APPLICANT

Company	General Medical Co.
Location	Mchenry, IL, US
510(k) history	15 submissions · 14 cleared · 1977-1993

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Device record: <https://www.510kdatabase.net/k802231/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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