

K780703 CUFF PACERJul 17, 1978
83 days to decisionK780703 · Product code: **CBA** · AnesthesiologySource: <https://www.510kdatabase.net/k780703/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Air Embolism, Ultrasonic (CBA)
Date received	Apr 25, 1978
Decision date	Jul 17, 1978
Days to decision	83 days
Third-party review	No

APPLICANT

Company	Omp Laboratories, Inc.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1976-1978

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Device record: <https://www.510kdatabase.net/k780703/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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