FDA 510(k) Premarket Notification

K780703 CUFF PACER

CLEARED

Jul 17, 1978
83 days to decision

K780703 · Product code: **CBA** · Anesthesiology
Source: 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