FDA 510(k) Premarket Notification

K833633 BRONCHITRAC DF DUAL RESP/CATHETER

CLEARED

Dec 8, 1983
52 days to decision

K833633 · Product code: **BSY** · Anesthesiology
Source: https://www.510kdatabase.net/k833633/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheters, Suction, Tracheobronchial (BSY)
Date received	Oct 17, 1983
Decision date	Dec 8, 1983
Days to decision	52 days
Third-party review	No

APPLICANT

Company	**Ackrad Laboratories**
Location	Mchenry, IL, US
510(k) history	42 submissions · 41 cleared · 1979-2002

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k833633/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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