FDA 510(k) Premarket Notification

K883899 MANUAL PERCUSSOR CUPS

CLEARED

Dec 13, 1988
89 days to decision

K883899 · Product code: **BYI** · Anesthesiology
Source: https://www.510kdatabase.net/k883899/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Percussor, Powered-electric (BYI)
Date received	Sep 15, 1988
Decision date	Dec 13, 1988
Days to decision	89 days
Third-party review	No

APPLICANT

Company	**Medical Components, Inc.**
Location	Mchenry, IL, US
Contact	DUNCAN G JOHNSON
510(k) history	63 submissions · 55 cleared · 1980-2025