FDA 510(k) Premarket Notification

K882604 PULMO-PATTY

CLEARED

Jan 4, 1989
194 days to decision

K882604 · Product code: **BYI** · Anesthesiology
Source: https://www.510kdatabase.net/k882604/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Percussor, Powered-electric (BYI)
Date received	Jun 24, 1988
Decision date	Jan 4, 1989
Days to decision	194 days
Third-party review	No

APPLICANT

Company	**Tower Stopper, Inc.**
Location	Hancock, MN, US
Contact	ARTHUR NOHL
510(k) history	1 submissions · 1 cleared · 1989-1989