FDA 510(k) Premarket Notification

K931988 3P (PULMONARY PERCUSSIVE PACK)

CLEARED

Oct 28, 1994
554 days to decision

K931988 · Product code: **BYI** · Anesthesiology
Source: https://www.510kdatabase.net/k931988/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Percussor, Powered-electric (BYI)
Date received	Apr 22, 1993
Decision date	Oct 28, 1994
Days to decision	554 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Delta Medical Manuf, Inc.**
Location	Raleigh, NC, US
Contact	DEN RAVELY
510(k) history	1 submissions · 1 cleared · 1994-1994