FDA 510(k) Premarket Notification

K924541 EPIDURAL MINIPACK

CLEARED

Jul 9, 1993
304 days to decision

K924541 · Product code: **BYO** · Anesthesiology
Source: https://www.510kdatabase.net/k924541/

SUBMISSION DETAILS

Decision	Substantially Equivalent - K
Submission type	Traditional
Device classification	Bottle, Blow (BYO)
Date received	Sep 8, 1992
Decision date	Jul 9, 1993
Days to decision	304 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Concord/Portex**
Location	Keene, NH, US
Contact	ROBERT WHEELER
510(k) history	23 submissions · 20 cleared · 1989-1993

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k924541/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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