FDA 510(k) Premarket Notification

K896121 SCHILLER MODEL SP-200 SPIROVIT(TM)

CLEARED

Jan 9, 1990
77 days to decision

K896121 · Product code: **BZG** · Anesthesiology
Source: https://www.510kdatabase.net/k896121/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Diagnostic (BZG)
Date received	Oct 24, 1989
Decision date	Jan 9, 1990
Days to decision	77 days
Third-party review	No

APPLICANT

Company	**Schiller America, Inc.**
Location	Tustin, CA, US
Contact	GODFREY J FLETCHER
510(k) history	10 submissions · 10 cleared · 1990-1998