FDA 510(k) Premarket Notification

K914272 KOKO SPIROMETER

CLEARED

Jul 2, 1992
282 days to decision

K914272 · Product code: **BZG** · Anesthesiology
Source: https://www.510kdatabase.net/k914272/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Diagnostic (BZG)
Date received	Sep 24, 1991
Decision date	Jul 2, 1992
Days to decision	282 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Ferraris Respiratory, Inc.**
Location	Louisville, CO, US
Contact	ARLIN LEHMAN
510(k) history	2 submissions · 2 cleared · 1992-2002