FDA 510(k) Premarket Notification

K020102 IQTEQ SPIROMETER

CLEARED

Mar 21, 2003
434 days to decision

K020102 · Product code: **BZG** · Anesthesiology
Source: https://www.510kdatabase.net/k020102/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Diagnostic (BZG)
Date received	Jan 11, 2002
Decision date	Mar 21, 2003
Days to decision	434 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Iqteq Development**
Location	Cape Town, Western Cape, ZA
Contact	RAY WRIGHT
510(k) history	1 submissions · 1 cleared · 2003-2003