FDA 510(k) Premarket Notification

K981725 OPTOVENT RESPIRATORY MONITOR MODEL NUMBER RR 9700

CLEARED

Feb 2, 1999
263 days to decision

K981725 · Product code: **BZQ** · Anesthesiology
Source: https://www.510kdatabase.net/k981725/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Breathing Frequency (BZQ)
Date received	May 15, 1998
Decision date	Feb 2, 1999
Days to decision	263 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Optovent, Inc.**
Location	Fort Lee, NJ, US
Contact	CLAES RYMOND
510(k) history	1 submissions · 1 cleared · 1999-1999