

**K981725 OPTOVENT RESPIRATORY MONITOR MODEL
NUMBER RR 9700**

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k981725/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 25, 2026