FDA 510(k) Premarket Notification

K820248 RODDER VOLUME VENTILATOR

CLEARED

Apr 16, 1982
77 days to decision

K820248 · Product code: **CBK** · Anesthesiology
Source: https://www.510kdatabase.net/k820248/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Continuous, Facility Use (CBK)
Date received	Jan 29, 1982
Decision date	Apr 16, 1982
Days to decision	77 days
Third-party review	No

APPLICANT

Company	**Pharmaquest Corp.**
Location	Walker, MI, US
510(k) history	14 submissions · 14 cleared · 1981-1995