FDA 510(k) Premarket Notification

K874198 VITALOG HMS-3000

CLEARED

Mar 22, 1988
165 days to decision

K874198 · Product code: **DQA** · Anesthesiology
Source: https://www.510kdatabase.net/k874198/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Oct 9, 1987
Decision date	Mar 22, 1988
Days to decision	165 days
Third-party review	No

APPLICANT

Company	**Vitalog Corp.**
Location	Walker, MI, US
Contact	BRUCE RULE
510(k) history	4 submissions · 4 cleared · 1981-1988