FDA 510(k) Premarket Notification

K902758 COBE SATURATION/HEMATOCRIT MONITOR W/ FIBER OPTIC

CLEARED

Aug 23, 1990
59 days to decision

K902758 · Product code: **DQA** · Anesthesiology
Source: https://www.510kdatabase.net/k902758/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Jun 25, 1990
Decision date	Aug 23, 1990
Days to decision	59 days
Third-party review	No

APPLICANT

Company	**Cobe Laboratories, Inc.**
Location	Mchenry, IL, US
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510(k) history	77 submissions · 77 cleared · 1976-1993