

**K902758 COBE SATURATION/HEMATOCRIT MONITOR W/
FIBER OPTIC**Aug 23, 1990
59 days to decisionK902758 · 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
Contact	MARY L ARMSTRONG
Website	https://www.gambro.com
510(k) history	77 submissions · 77 cleared · 1976-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k902758/>; 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 26, 2026