

K911207 OXICOMPLUSAug 26, 1991
160 days to decisionK911207 · Product code: **DQA** · AnesthesiologySource: <https://www.510kdatabase.net/k911207/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Mar 19, 1991
Decision date	Aug 26, 1991
Days to decision	160 days
Third-party review	No

APPLICANT

Company	Waters Instruments, Inc.
Location	Walker, MI, US
Contact	STEVE BOUTELLE
510(k) history	12 submissions · 12 cleared · 1981-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k911207/>; 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