

K880448 CERION OXY THRESHOLD ALARMMar 21, 1988
48 days to decisionK880448 · Product code: **CCL** · AnesthesiologySource: <https://www.510kdatabase.net/k880448/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Oxygen, Gaseous-phase (CCL)
Date received	Feb 2, 1988
Decision date	Mar 21, 1988
Days to decision	48 days
Third-party review	No

APPLICANT

Company	Ceramatec, Inc.
Location	Salt Lake City, UT, US
Contact	MARK HASLAM
510(k) history	4 submissions · 4 cleared · 1988-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k880448/>; 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 25, 2026