

K882273 HORIZON 2000 PULSE OXIMETRY AND MODULEAug 25, 1988
86 days to decisionK882273 · Product code: **DQA** · AnesthesiologySource: <https://www.510kdatabase.net/k882273/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	May 31, 1988
Decision date	Aug 25, 1988
Days to decision	86 days
Third-party review	No

APPLICANT

Company	Mennen Medical, Inc.
Location	Clarence, NY, US
Contact	THOMAS W CONNELLY
510(k) history	34 submissions · 34 cleared · 1985-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k882273/> 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 21, 2026