

K895987 KELLER PULSE OXIMETER - KMS 850Feb 22, 1990
133 days to decisionK895987 · Product code: **DQA** · AnesthesiologySource: <https://www.510kdatabase.net/k895987/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Oct 12, 1989
Decision date	Feb 22, 1990
Days to decision	133 days
Third-party review	No

APPLICANT

Company	Keller Medical Specialties Products, Inc.
Location	Antioch, IL, US
Contact	JAY KELLER
510(k) history	6 submissions · 6 cleared · 1990-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k895987/>; 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