

K871864 PB240 PATIENT MONITOR/PULSE OXIMETERJun 24, 1987
42 days to decisionK871864 · Product code: **DQA** · AnesthesiologySource: <https://www.510kdatabase.net/k871864/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	May 13, 1987
Decision date	Jun 24, 1987
Days to decision	42 days
Third-party review	No

APPLICANT

Company	Puritan Bennett Corp.
Location	Mchenry, IL, US
Contact	HARTLEY C ERICSON
510(k) history	110 submissions · 101 cleared · 1976-2007

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k871864/>; 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