FDA 510(k) Premarket Notification

K882273 HORIZON 2000 PULSE OXIMETRY AND MODULE

CLEARED

Aug 25, 1988
86 days to decision

K882273 · Product code: **DQA** · Anesthesiology
Source: 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