FDA 510(k) Premarket Notification

K904039 REFLECTANCE SENSOR (RS-10)

CLEARED

Dec 27, 1990
114 days to decision

K904039 · Product code: **DQA** · Anesthesiology
Source: https://www.510kdatabase.net/k904039/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Sep 4, 1990
Decision date	Dec 27, 1990
Days to decision	114 days
Third-party review	No

APPLICANT

Company	**Nellcor, Inc.**
Location	Mchenry, IL, US
Contact	KENNETH MICHAEL
510(k) history	18 submissions · 18 cleared · 1982-1995