FDA 510(k) Premarket Notification

K811391 OXIMETRIX ACCU SAT CPB

CLEARED

Jul 20, 1981
63 days to decision

K811391 · Product code: **DRY** · Cardiovascular
Source: https://www.510kdatabase.net/k811391/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Blood-gas, On-line, Cardiopulmonary Bypass (DRY)
Date received	May 18, 1981
Decision date	Jul 20, 1981
Days to decision	63 days
Third-party review	No

APPLICANT

Company	**Oximetrix, Inc.**
Location	Mchenry, IL, US
510(k) history	12 submissions · 12 cleared · 1976-1986