FDA 510(k) Premarket Notification

K792081 SPECTRA CENTRAL PT MONITORING SYS

CLEARED

Nov 13, 1979
28 days to decision

K792081 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k792081/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Oct 16, 1979
Decision date	Nov 13, 1979
Days to decision	28 days
Third-party review	No

APPLICANT

Company	**American Optical Corp.**
Location	Mchenry, IL, US
510(k) history	35 submissions · 35 cleared · 1976-1995