FDA 510(k) Premarket Notification

K781845 CAMSCAN SYSTEM

CLEARED

Nov 20, 1978
19 days to decision

K781845 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k781845/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Nov 1, 1978
Decision date	Nov 20, 1978
Days to decision	19 days
Third-party review	No

APPLICANT

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