FDA 510(k) Premarket Notification

K802179 CARDIOSCOPE J200

CLEARED

Nov 12, 1980
64 days to decision

K802179 · Product code: **DRS** · Cardiovascular
Source: https://www.510kdatabase.net/k802179/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transducer, Blood-pressure, Extravascular (DRS)
Date received	Sep 9, 1980
Decision date	Nov 12, 1980
Days to decision	64 days
Third-party review	No

APPLICANT

Company	**Telectronics, Inc.**
Location	Mchenry, IL, US
510(k) history	107 submissions · 107 cleared · 1977-1990