FDA 510(k) Premarket Notification

K810588 SYNCHRONIZER #208

CLEARED

Apr 10, 1981
37 days to decision

K810588 · Product code: **IXO** · Radiology
Source: https://www.510kdatabase.net/k810588/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Synchronizer, Ecg / Respirator, Radiographic (IXO)
Date received	Mar 4, 1981
Decision date	Apr 10, 1981
Days to decision	37 days
Third-party review	No

APPLICANT

Company	**Brattle Instrument Corp.**
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1977-1981