FDA 510(k) Premarket Notification

K791632 COMPU-SCAN

CLEARED

Nov 13, 1979
84 days to decision

K791632 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k791632/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Aug 21, 1979
Decision date	Nov 13, 1979
Days to decision	84 days
Third-party review	No

APPLICANT

Company	**Hittman Medical Systems**
Location	Mchenry, IL, US
510(k) history	3 submissions · 3 cleared · 1979-1983