FDA 510(k) Premarket Notification

K792105 CRITIKON DUAL CHANNEL RECORDER

CLEARED

Oct 26, 1979
4 days to decision

K792105 · Product code: **DSF** · Cardiovascular
Source: https://www.510kdatabase.net/k792105/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Paper Chart (DSF)
Date received	Oct 22, 1979
Decision date	Oct 26, 1979
Days to decision	4 days
Third-party review	No

APPLICANT

Company	**Critikon Company, LLC**
Location	Mchenry, IL, US
510(k) history	51 submissions · 51 cleared · 1979-2000