FDA 510(k) Premarket Notification

K761023 SIMULATOR, 10 LEAD ECG

CLEARED

Jan 31, 1977
81 days to decision

K761023 · Product code: **DPS** · Cardiovascular
Source: https://www.510kdatabase.net/k761023/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrocardiograph (DPS)
Date received	Nov 11, 1976
Decision date	Jan 31, 1977
Days to decision	81 days
Third-party review	No

APPLICANT

Company	**Birtcher Corp.**
Location	Mchenry, IL, US
510(k) history	27 submissions · 27 cleared · 1976-1988