FDA 510(k) Premarket Notification

K770330 MONITOR, PULSE WAVE VELOCITY P607

CLEARED

Feb 24, 1977
7 days to decision

K770330 · Product code: **DXG** · Cardiovascular
Source: https://www.510kdatabase.net/k770330/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Pre-programmed, Single-function (DXG)
Date received	Feb 17, 1977
Decision date	Feb 24, 1977
Days to decision	7 days
Third-party review	No

APPLICANT

Company	**Cyborg Corp.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1977-1977