FDA 510(k) Premarket Notification

K802576 EXTERNAL JUGULAR CVP CATHERIZATION KIT

CLEARED

Nov 12, 1980
23 days to decision

K802576 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k802576/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Oct 20, 1980
Decision date	Nov 12, 1980
Days to decision	23 days
Third-party review	No

APPLICANT

Company	**Willson**
Location	Walker, MI, US
510(k) history	2 submissions · 2 cleared · 1980-1981