FDA 510(k) Premarket Notification

K832516 MAGNATRODE CARDIAC MONITORING ELECTRODE

CLEARED

Dec 16, 1983
141 days to decision

K832516 · Product code: **DRX** · Cardiovascular
Source: https://www.510kdatabase.net/k832516/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jul 28, 1983
Decision date	Dec 16, 1983
Days to decision	141 days
Third-party review	No

APPLICANT

Company	**R2 Corp.**
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1980-1983