

K800935 R2 APEX-POST.,ANT.&POST. ELECT.(12 MODS)May 23, 1980
32 days to decisionK800935 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k800935/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Apr 21, 1980
Decision date	May 23, 1980
Days to decision	32 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k800935/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 1, 2026