

K802576 EXTERNAL JUGULAR CVP CATHERIZATION KITNov 12, 1980
23 days to decisionK802576 · Product code: **DRX** · CardiovascularSource: <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

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Device record: <https://www.510kdatabase.net/k802576/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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