

K810642 ATRIAL-VENTRICULAR PACING ELECTRODE KITApr 8, 1981
29 days to decisionK810642 · Product code: **LDF** · CardiovascularSource: <https://www.510kdatabase.net/k810642/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Mar 10, 1981
Decision date	Apr 8, 1981
Days to decision	29 days
Third-party review	No

APPLICANT

Company	Stanco Medical, Inc.
Location	Mchenry, IL, US
510(k) history	19 submissions · 19 cleared · 1979-1981

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Device record: <https://www.510kdatabase.net/k810642/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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