

K843141 CRITIKON HEPARIN-COATED CARDIOVASCULARSep 4, 1984
26 days to decisionK843141 · Product code: **LDF** · CardiovascularSource: <https://www.510kdatabase.net/k843141/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Aug 9, 1984
Decision date	Sep 4, 1984
Days to decision	26 days
Third-party review	No

APPLICANT

Company	Critikon Company, LLC
Location	Mchenry, IL, US
510(k) history	51 submissions · 51 cleared · 1979-2000

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Device record: <https://www.510kdatabase.net/k843141/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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