

**K792105 CRITIKON DUAL CHANNEL RECORDER**Oct 26, 1979  
4 days to decisionK792105 · Product code: **DSF** · CardiovascularSource: <https://www.510kdatabase.net/k792105/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Paper Chart (DSF)
Date received	Oct 22, 1979
Decision date	Oct 26, 1979
Days to decision	4 days
Third-party review	No

**APPLICANT**

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

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k792105/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 26, 2026