

**K930196 CEPTI-SEAL I.V. PREP KIT**Jul 2, 1993  
169 days to decisionK930196 · Product code: **LRS** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k930196/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - KD
Submission type	Traditional
Device classification	I.v. Start Kit (LRS)
Date received	Jan 14, 1993
Decision date	Jul 2, 1993
Days to decision	169 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Medi-Flex Hospital Products, Inc.</b>
Location	Overland Park, KS, US
Contact	ORLANDO CORDOVA
510(k) history	13 submissions · 9 cleared · 1991-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k930196/> 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