

**K914475 IV START KIT**Mar 18, 1992  
162 days to decisionK914475 · Product code: **LRS** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k914475/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - KD
Submission type	Traditional
Device classification	I.v. Start Kit (LRS)
Date received	Oct 8, 1991
Decision date	Mar 18, 1992
Days to decision	162 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Cypress Medical Products, Ltd.</b>
Location	Mc Henry, IL, US
Contact	VARUN SONI
510(k) history	26 submissions · 19 cleared · 1989-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k914475/> 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 June 29, 2026