

**K945160 PARACENTESIS NEEDLES/CANNULAS**Nov 14, 1994  
24 days to decisionK945160 · Product code: **LRO** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k945160/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	General Surgery Tray (LRO)
Date received	Oct 21, 1994
Decision date	Nov 14, 1994
Days to decision	24 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Tri-Med Specialties, Inc.</b>
Location	Overland Park, KS, US
Contact	MIKE KNOTH
510(k) history	14 submissions · 12 cleared · 1985-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k945160/> 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 July 1, 2026