

**K012042 KINETIC CANNULA**Sep 20, 2001  
83 days to decisionK012042 · Product code: **QPB** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k012042/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Suction, Lipoplasty For Removal (QPB)
Date received	Jun 29, 2001
Decision date	Sep 20, 2001
Days to decision	83 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Reliance Medical Corp.</b>
Location	Tucson, AZ, US
Contact	ROBERT W NICKS
510(k) history	3 submissions · 3 cleared · 1985-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k012042/> 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 3, 2026