

**K890170 SCISSORS**Feb 3, 1989  
25 days to decisionK890170 · Product code: **LRW** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k890170/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Scissors, General, Surgical (LRW)
Date received	Jan 9, 1989
Decision date	Feb 3, 1989
Days to decision	25 days
Third-party review	No

**APPLICANT**

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Company	<b>Dhason Brothers, Inc.</b>
Location	Hyattsville, MD, US
Contact	ROBERT DHASON
510(k) history	14 submissions · 14 cleared · 1989-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890170/>, 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 22, 2026