

**K910823 PERCUDISC(TM) SYSTEM**Dec 30, 1991  
307 days to decisionK910823 · Product code: **GEX** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k910823/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Powered Laser Surgical Instrument (GEX)
Date received	Feb 26, 1991
Decision date	Dec 30, 1991
Days to decision	307 days
Third-party review	No

**APPLICANT**

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Company	<b>Percudisc, Inc.</b>
Location	Mountain View, CA, US
Contact	CHARLES L ROSE
510(k) history	2 submissions · 1 cleared · 1991-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k910823/> 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 7, 2026