

**K833021 PERI-GUARD PERICARDIAL PATCH-HERNIA**Dec 26, 1983  
111 days to decisionK833021 · Product code: **DXZ** · CardiovascularSource: <https://www.510kdatabase.net/k833021/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Patch, Pledget And Intracardiac, Petp, Ptfе, Polypropylene (DXZ)
Date received	Sep 6, 1983
Decision date	Dec 26, 1983
Days to decision	111 days
Third-party review	No

**APPLICANT**

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Company	<b>Genetic</b>
Location	Mchenry, IL, US
Website	<a href="http://www.myriad.com">http://www.myriad.com</a>
510(k) history	1 submissions · 1 cleared · 1983-1983

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k833021/>, 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 8, 2026