

**K963437 PELVIC HOLDERS**Nov 19, 1996  
81 days to decisionK963437 · Product code: **FMQ** · General Hospital  
Source: <https://www.510kdatabase.net/k963437/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Restraint, Protective (FMQ)
Date received	Aug 30, 1996
Decision date	Nov 19, 1996
Days to decision	81 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>J. T. Posey Co.</b>
Location	Arcadia, CA, US
Contact	MICHAEL KEEFE
510(k) history	13 submissions · 13 cleared · 1992-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k963437/> 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 June 14, 2026