FDA 510(k) Premarket Notification

K963437 PELVIC HOLDERS

CLEARED

Nov 19, 1996
81 days to decision

K963437 · Product code: **FMQ** · General Hospital
Source: https://www.510kdatabase.net/k963437/

SUBMISSION DETAILS

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

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