FDA 510(k) Premarket Notification

K962429 SAFETY VEST

CLEARED

Sep 18, 1996
86 days to decision

K962429 · Product code: **FMQ** · General Hospital
Source: https://www.510kdatabase.net/k962429/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Restraint, Protective (FMQ)
Date received	Jun 24, 1996
Decision date	Sep 18, 1996
Days to decision	86 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Johnson & Johnson Professionals, Inc.**
Location	Raynham, MA, US
Contact	JOHANNA ADLER
Website	https://www.jnj.com
510(k) history	206 submissions · 184 cleared · 1976-2000

Johnson & Johnson Professionals, Inc. is a medical device company based in Raynham, Massachusetts. The company specializes in surgical and orthopedic devices. The company has received FDA 510(k) clearances from total submissions between 1976 and 2000. Orthopedic devices and neurosurgical instruments represent core product categories. Notable cleared devices include hip and elbow prostheses, programmable valve systems, and aneurysm clips. The company is inactive and represents a historical regulatory record with no submissions in more than two decades. Explore the complete...