FDA 510(k) Premarket Notification

K760868 VEST RESTRAINT

CLEARED

Nov 1, 1976
14 days to decision

K760868 · Product code: **KNW** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k760868/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Instrument, Biopsy (KNW)
Date received	Oct 18, 1976
Decision date	Nov 1, 1976
Days to decision	14 days
Third-party review	No

APPLICANT

Company	**Orthopedic Equipment Co., Inc.**
Location	Mchenry, IL, US
510(k) history	45 submissions · 45 cleared · 1976-1984