FDA 510(k) Premarket Notification

K760618 PATIENT RESTRAINT

CLEARED

Nov 24, 1976
72 days to decision

K760618 · Product code: **FMQ** · General Hospital
Source: https://www.510kdatabase.net/k760618/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Restraint, Protective (FMQ)
Date received	Sep 13, 1976
Decision date	Nov 24, 1976
Days to decision	72 days
Third-party review	No

APPLICANT

Company	**Encompass Group, LLC**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1976-2020