

K760618 PATIENT RESTRAINTNov 24, 1976
72 days to decisionK760618 · Product code: **FMQ** · General HospitalSource: <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

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Device record: <https://www.510kdatabase.net/k760618/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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