FDA 510(k) Premarket Notification

K962327 SEKURE SEAT ADULT PASSIVE RESTRAINING DEVICE

CLEARED

Mar 6, 1997
262 days to decision

K962327 · Product code: **FMQ** · General Hospital
Source: https://www.510kdatabase.net/k962327/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Restraint, Protective (FMQ)
Date received	Jun 17, 1996
Decision date	Mar 6, 1997
Days to decision	262 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Sekure Seat Corp.**
Location	Hickory, NC, US
Contact	LAURA ANN REID
510(k) history	1 submissions · 1 cleared · 1997-1997