

**K962327 SEKURE SEAT ADULT PASSIVE RESTRAINING  
DEVICE**Mar 6, 1997  
262 days to decisionK962327 · Product code: **FMQ** · General Hospital  
Source: <https://www.510kdatabase.net/k962327/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k962327/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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