FDA 510(k) Premarket Notification

K953735 AUTOJECT MINI

CLEARED

Nov 8, 1995
90 days to decision

K953735 · Product code: **KZH** · General Hospital
Source: https://www.510kdatabase.net/k953735/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Introducer, Syringe Needle (KZH)
Date received	Aug 10, 1995
Decision date	Nov 8, 1995
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Owen Mumford USA, Inc.**
Location	Marietta, GA, US
Contact	ROBERT SHAW
510(k) history	10 submissions · 10 cleared · 1995-2001

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k953735/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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