FDA 510(k) Premarket Notification

K952610 PLEXUS/3000 (P/3000), PLEXUS/3500 (P/3500)

CLEARED

Sep 13, 1995
98 days to decision

K952610 · Product code: **FNM** · General Hospital
Source: https://www.510kdatabase.net/k952610/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mattress, Air Flotation, Alternating Pressure (FNM)
Date received	Jun 7, 1995
Decision date	Sep 13, 1995
Days to decision	98 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Plexus Medical**
Location	Covina, CA, US
Contact	DAVBID BUCHICCHIO
510(k) history	3 submissions · 3 cleared · 1994-1995