

**K952610 PLEXUS/3000 (P/3000), PLEXUS/3500 (P/3500)**Sep 13, 1995  
98 days to decisionK952610 · Product code: **FNM** · General Hospital  
Source: <https://www.510kdatabase.net/k952610/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k952610/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 6, 2026