

**K961854 SPR PLUS II OVERLAY SYSTEM (CL250/CL212)**Aug 9, 1996  
87 days to decisionK961854 · Product code: **FNM** · General Hospital  
Source: <https://www.510kdatabase.net/k961854/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mattress, Air Flotation, Alternating Pressure (FNM)
Date received	May 14, 1996
Decision date	Aug 9, 1996
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Gaymar Industries, Inc.</b>
Location	Mchenry, IL, US
Contact	PETER SCOTT
Website	<a href="https://www.gaymar.com">https://www.gaymar.com</a>
510(k) history	27 submissions · 27 cleared · 1976-2011

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k961854/> 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 June 20, 2026