

**K910830 INVALIDIFT**Jun 9, 1992  
468 days to decisionK910830 · Product code: **FRZ** · General Hospital  
Source: <https://www.510kdatabase.net/k910830/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Patient Transfer, Powered (FRZ)
Date received	Feb 27, 1991
Decision date	Jun 9, 1992
Days to decision	468 days
Third-party review	No

**APPLICANT**

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Company	<b>American Invalift Corp.</b>
Location	White Plains, NY, US
Contact	THERESE RUBINO
510(k) history	1 submissions · 1 cleared · 1992-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k910830/>, 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 7, 2026