

**K905756 TITAN MERIDIAN INTERSEGMENTAL ROLLERTABLE**Jan 8, 1991  
12 days to decisionK905756 · Product code: **JFB** · Physical MedicineSource: <https://www.510kdatabase.net/k905756/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Physical Therapy, Multi Function (JFB)
Date received	Dec 27, 1990
Decision date	Jan 8, 1991
Days to decision	12 days
Third-party review	No

**APPLICANT**

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Company	<b>Titan Technology Intl., Inc.</b>
Location	St. Louis, MI, US
Contact	HOWARD COOPER
510(k) history	1 submissions · 1 cleared · 1991-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k905756/>; 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 30, 2026