

**K810220 POWERED PATIENT SYSTEM**Feb 4, 1981  
8 days to decisionK810220 · Product code: **IKZ** · Physical MedicineSource: <https://www.510kdatabase.net/k810220/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bed, Patient Rotation, Powered (IKZ)
Date received	Jan 27, 1981
Decision date	Feb 4, 1981
Days to decision	8 days
Third-party review	No

**APPLICANT**

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Company	<b>Tri W-G, Inc.</b>
Location	Mchenry, IL, US
510(k) history	35 submissions · 35 cleared · 1979-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k810220/> 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 May 21, 2026