

**K905113 ROTO REST**Nov 26, 1990  
13 days to decisionK905113 · Product code: **IKZ** · Physical MedicineSource: <https://www.510kdatabase.net/k905113/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bed, Patient Rotation, Powered (IKZ)
Date received	Nov 13, 1990
Decision date	Nov 26, 1990
Days to decision	13 days
Third-party review	No
Other names	TRAUMA AND INTENSIVE CARE BED

**APPLICANT**

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Company	<b>Ethos Medical Products , Ltd.</b>
Location	Athlone, IE
Contact	PAT CONNOLLY
510(k) history	1 submissions · 1 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k905113/> 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