

K962788 CLINIDYNE ROTATIONAL THERAPY MATTRESS SYSTEMApr 30, 1997
287 days to decisionK962788 · Product code: **IKZ** · Physical Medicine
Source: <https://www.510kdatabase.net/k962788/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bed, Patient Rotation, Powered (IKZ)
Date received	Jul 17, 1996
Decision date	Apr 30, 1997
Days to decision	287 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k962788/> Data sourced from FDA 510(k) public records (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. - Generated June 20, 2026