

K010252 MAGNUMApr 26, 2001
90 days to decisionK010252 · Product code: **JFB** · Physical Medicine
Source: <https://www.510kdatabase.net/k010252/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Physical Therapy, Multi Function (JFB)
Date received	Jan 26, 2001
Decision date	Apr 26, 2001
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Magnum Equipment LLC
Location	Oklahoma City, OK, US
Contact	MONTE REGAL
510(k) history	1 submissions · 1 cleared · 2001-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k010252/> 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 29, 2026