FDA 510(k) Premarket Notification

K010252 MAGNUM

CLEARED

Apr 26, 2001
90 days to decision

K010252 · 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