510(k) Premarket Notification

1 to 4 of 4 Results
Applicant: *axiom worldwide* **Decision Date**
To: *01/15/2020*

Results per Page 10

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Device Name	Applicant	510(K) Number	Decision Date
Drx9000 True Spinal Decompression System	AXIOM WORLDWIDE, INC.	K060735	05/26/2006
Axiom Nvp8500	AXIOM WORLDWIDE, INC.	K051135	08/11/2005
Axiom Eps8000	AXIOM WORLDWIDE, INC.	K050687	07/28/2005
Drx 2000	AXIOM WORLDWIDE, INC.	K010292	05/01/2001

Page Last Updated: 01/13/2020
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