FDA 510(k) Premarket Notification

K032456 PROWESS

CLEARED

Oct 16, 2003
66 days to decision

K032456 · Product code: **MUJ** · Radiology
Source: https://www.510kdatabase.net/k032456/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Planning, Radiation Therapy Treatment (MUJ)
Date received	Aug 11, 2003
Decision date	Oct 16, 2003
Days to decision	66 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Prowess, Inc.**
Location	Chico, CA, US
Contact	W. JAMES BISHOP
510(k) history	9 submissions · 9 cleared · 2003-2023