FDA 510(k) Premarket Notification

K060349 PROGUIDE NEEDLE SET

CLEARED

Mar 10, 2006
28 days to decision

K060349 · Product code: **JAQ** · Radiology
Source: https://www.510kdatabase.net/k060349/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Applicator, Radionuclide, Remote-controlled (JAQ)
Date received	Feb 10, 2006
Decision date	Mar 10, 2006
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Nucletron Corporation**
Location	Columbia, MD, US
Contact	LISA DIMMICK
510(k) history	19 submissions · 19 cleared · 2002-2011