

K060349 PROGUIDE NEEDLE SETMar 10, 2006
28 days to decisionK060349 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k060349/> 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 May 17, 2026