FDA 510(k) Premarket Notification

CLEARED

K080934 VIENNA RING CT/MR APPLICATOR SET, ROUND POINT TITANIUM NEEDLE SET, TROCAR POINT TITANIUM NEEDLE SET

May 28, 2008
56 days to decision

K080934 · Product code: **JAQ** · Radiology
Source: https://www.510kdatabase.net/k080934/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Applicator, Radionuclide, Remote-controlled (JAQ)
Date received	Apr 2, 2008
Decision date	May 28, 2008
Days to decision	56 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