

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k080934/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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