FDA 510(k) Premarket Notification

K041782 SEEMOR 5.0 IMAGE DISPLAY PROGRAM

CLEARED

Aug 16, 2004
46 days to decision

K041782 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k041782/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Jul 1, 2004
Decision date	Aug 16, 2004
Days to decision	46 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Areeda Assoc., Ltd.**
Location	Los Angeles, CA, US
Contact	JOSEPH AREEDA
510(k) history	2 submissions · 2 cleared · 1997-2004