

**K041782 SEEMOR 5.0 IMAGE DISPLAY PROGRAM**Aug 16, 2004  
46 days to decisionK041782 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k041782/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k041782/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 18, 2026