

**K000770 PROVIEW MEDICAL IMAGE DISPLAY MODULE**May 16, 2000  
68 days to decisionK000770 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k000770/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Mar 9, 2000
Decision date	May 16, 2000
Days to decision	68 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Cerner Corp.</b>
Location	Kansas City, MO, US
Contact	SHELLEY LOOBY
510(k) history	7 submissions · 7 cleared · 1996-2012

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k000770/> 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 17, 2026