FDA 510(k) Premarket Notification

K000770 PROVIEW MEDICAL IMAGE DISPLAY MODULE

CLEARED

May 16, 2000
68 days to decision

K000770 · 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	**Cerner Corp.**
Location	Kansas City, MO, US
Contact	SHELLEY LOOBY
510(k) history	7 submissions · 7 cleared · 1996-2012