FDA 510(k) Premarket Notification

K071492 FIRSTVIEW

CLEARED

Aug 28, 2007
90 days to decision

K071492 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k071492/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	May 30, 2007
Decision date	Aug 28, 2007
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Riverain Medical Group,Llc**
Location	Miamisburg, OH, US
Contact	JENNIFER STEINKE
510(k) history	2 submissions · 2 cleared · 2007-2011