FDA 510(k) Premarket Notification

K101842 ACIES

CLEARED

Feb 4, 2011
218 days to decision

K101842 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k101842/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Jul 1, 2010
Decision date	Feb 4, 2011
Days to decision	218 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Konica Minolta Medical & Graphic, Inc.**
Location	Hachioji-Shi Tokyo, JP
Contact	RUSSELL MUNVES
510(k) history	27 submissions · 27 cleared · 2002-2013