FDA 510(k) Premarket Notification

K082135 SHOWCASE

CLEARED

Oct 21, 2008
84 days to decision

K082135 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k082135/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Jul 29, 2008
Decision date	Oct 21, 2008
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Trillium Technology, Inc.**
Location	San Leandro, CA, US
Contact	GARY J ALLSEBROOK
510(k) history	1 submissions · 1 cleared · 2008-2008