FDA 510(k) Premarket Notification

K082347 DIGITAL RADIOGRAPHY, FLEXDR C30

CLEARED

Nov 18, 2008
95 days to decision

K082347 · Product code: **MQB** · Radiology
Source: https://www.510kdatabase.net/k082347/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Solid State X-ray Imager (flat Panel/digital Imager) (MQB)
Date received	Aug 15, 2008
Decision date	Nov 18, 2008
Days to decision	95 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Konica Minolta Technoproduct Co., Ltd.**
Location	New York, NY, US
Contact	RUSSEL MUNVES
510(k) history	1 submissions · 1 cleared · 2008-2008