FDA 510(k) Premarket Notification

K120484 VISIA ONCOLOGY

CLEARED

Mar 27, 2012
39 days to decision

K120484 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k120484/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Feb 17, 2012
Decision date	Mar 27, 2012
Days to decision	39 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Mevis Medical Solutions AG**
Location	Pewaukee, WI, US
Contact	THOMAS E TYNES
510(k) history	6 submissions · 6 cleared · 2011-2023