FDA 510(k) Premarket Notification

K070205 ZIOSTATION

CLEARED

Mar 23, 2007
60 days to decision

K070205 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k070205/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Jan 22, 2007
Decision date	Mar 23, 2007
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Ziosoft, Inc.**
Location	Irvine, CA, US
Contact	MARC GOODMAN
510(k) history	6 submissions · 6 cleared · 2007-2020