

K070205 ZIOSTATIONMar 23, 2007
60 days to decisionK070205 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k070205/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 19, 2026