FDA 510(k) Premarket Notification

K052379 MIM 3.5 (CIRCA)

CLEARED

Oct 31, 2005
62 days to decision

K052379 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k052379/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Aug 30, 2005
Decision date	Oct 31, 2005
Days to decision	62 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Mimvista Corp.**
Location	Cleveland, OH, US
Contact	PETER SIMMELINK
510(k) history	4 submissions · 4 cleared · 2005-2007