FDA 510(k) Premarket Notification

K974755 DYNA VUE 3-D IMAGING PRODUCT

CLEARED

Mar 4, 1998
75 days to decision

K974755 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k974755/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Dec 19, 1997
Decision date	Mar 4, 1998
Days to decision	75 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Imd Systems, Inc.**
Location	Amherst, NH, US
Contact	KENNETH M NICOLL
510(k) history	1 submissions · 1 cleared · 1998-1998