FDA 510(k) Premarket Notification

K955708 COMPUMED DICON CLIENT

CLEARED

Mar 1, 1996
78 days to decision

K955708 · Product code: **LMD** · Radiology
Source: https://www.510kdatabase.net/k955708/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Digital Image Communications, Radiological (LMD)
Date received	Dec 14, 1995
Decision date	Mar 1, 1996
Days to decision	78 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Compumed, Inc.**
Location	Mchenry, IL, US
Website	http://compumed.ning.com/
510(k) history	6 submissions · 6 cleared · 1977-1999