

K955708 COMPUMED DICON CLIENTMar 1, 1996
78 days to decisionK955708 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k955708/> 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 May 19, 2026