FDA 510(k) Premarket Notification

K992698 STENTOR, PAGEVIEW

CLEARED

Oct 25, 1999
75 days to decision

K992698 · Product code: **LMD** · Radiology
Source: https://www.510kdatabase.net/k992698/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Digital Image Communications, Radiological (LMD)
Date received	Aug 11, 1999
Decision date	Oct 25, 1999
Days to decision	75 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Stentor, Inc.**
Location	San Leandro, CA, US
Contact	GARY J ALLSEBROOK
510(k) history	3 submissions · 3 cleared · 1999-2004

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k992698/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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