

K993598 MAMMOGRAPHY PRO, MODEL 13017, DIAGNOSTIC PRO W/ MAMMOGRAPHY OPTION, MODEL 13205

Jan 20, 2000
87 days to decision

K993598 · Product code: **LMA** · Radiology
Source: <https://www.510kdatabase.net/k993598/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Digitizer, Image, Radiological (LMA)
Date received	Oct 25, 1999
Decision date	Jan 20, 2000
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Vidar Systems Corp.
Location	Herndon, VA, US
Contact	MARY "PENNIE" DRINKARD
510(k) history	13 submissions · 13 cleared · 1993-2013

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Device record: <https://www.510kdatabase.net/k993598/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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