

K884323 XIMAVIEW(TM)Apr 3, 1989
171 days to decisionK884323 · Product code: **KPQ** · Radiology
Source: <https://www.510kdatabase.net/k884323/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Simulation, Radiation Therapy (KPQ)
Date received	Oct 14, 1988
Decision date	Apr 3, 1989
Days to decision	171 days
Third-party review	No

APPLICANT

Company	Varian Assoc., Inc.
Location	Mchenry, IL, US
Contact	MILTON D SIEGEL
510(k) history	86 submissions · 86 cleared · 1976-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k884323/>; 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