

K912341 MULTICAM I 8X10Aug 26, 1991
94 days to decisionK912341 · Product code: **JAA** · Radiology
Source: <https://www.510kdatabase.net/k912341/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Fluoroscopic, Image-intensified (JAA)
Date received	May 24, 1991
Decision date	Aug 26, 1991
Days to decision	94 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Camtronics, Ltd.
Location	Delafield, WI, US
Contact	SCOTT J PEASE
510(k) history	16 submissions · 16 cleared · 1987-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k912341/> 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