FDA 510(k) Premarket Notification

K931974 AZ 92

CLEARED

Oct 8, 1993
169 days to decision

K931974 · Product code: **IWE** · Radiology
Source: https://www.510kdatabase.net/k931974/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Patient Position, Light-beam (IWE)
Date received	Apr 22, 1993
Decision date	Oct 8, 1993
Days to decision	169 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**A2j, Inc.**
Location	Houston, TX, US
Contact	JIM L.
510(k) history	4 submissions · 4 cleared · 1988-1993