FDA 510(k) Premarket Notification

K991435 AII-PRO 100 PLUS AND ALL-PRO 200

CLEARED

May 20, 1999
24 days to decision

K991435 · Product code: **IXW** · Radiology
Source: https://www.510kdatabase.net/k991435/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Processor, Radiographic-film, Automatic (IXW)
Date received	Apr 26, 1999
Decision date	May 20, 1999
Days to decision	24 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**All-Pro Imaging Corp.**
Location	Hicksville, NY, US
Contact	FREDERICK R FISCHER
510(k) history	4 submissions · 4 cleared · 1991-2000