FDA 510(k) Premarket Notification

K952188 ADVIEWER 2000

CLEARED

May 30, 1995
21 days to decision

K952188 · Product code: **IXC** · Radiology
Source: https://www.510kdatabase.net/k952188/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Illuminator, Radiographic-film (IXC)
Date received	May 9, 1995
Decision date	May 30, 1995
Days to decision	21 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Adp , Ltd.**
Location	Haifa, IL
Contact	DAN BEN-ZEEV
510(k) history	1 submissions · 1 cleared · 1995-1995

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k952188/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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