FDA 510(k) Premarket Notification

CLEARED

K926083 NUCLEAR PATIENT SANNING TABLES, VARIOUS TYPES

Feb 12, 1993
73 days to decision

K926083 · Product code: **IYX** · Radiology
Source: https://www.510kdatabase.net/k926083/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Camera, Scintillation (gamma) (IYX)
Date received	Dec 1, 1992
Decision date	Feb 12, 1993
Days to decision	73 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Biodan Medical Systems, Ltd.**
Location	Israel, IL
Contact	CLYDE SCHLEIN
510(k) history	20 submissions · 20 cleared · 1982-1993