

K926083 NUCLEAR PATIENT SANNING TABLES, VARIOUS TYPESFeb 12, 1993
73 days to decisionK926083 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k926083/> 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 20, 2026