FDA 510(k) Premarket Notification

K925705 ACCURAD-200 SYSTEM & ACCURAD-300 PLUS SYSTEM

CLEARED

Dec 28, 1992
46 days to decision

K925705 · Product code: **IWY** · Radiology
Source: https://www.510kdatabase.net/k925705/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Holder, Head, Radiographic (IWY)
Date received	Nov 12, 1992
Decision date	Dec 28, 1992
Days to decision	46 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Denar Corp.**
Location	Anaheim, CA, US
Contact	THOMAS J MULDOON
510(k) history	5 submissions · 5 cleared · 1985-1992