FDA 510(k) Premarket Notification

K890066 STAR 36

CLEARED

Mar 28, 1989
78 days to decision

K890066 · Product code: **IZO** · Radiology
Source: https://www.510kdatabase.net/k890066/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, High-voltage, X-ray, Diagnostic (IZO)
Date received	Jan 9, 1989
Decision date	Mar 28, 1989
Days to decision	78 days
Third-party review	No

APPLICANT

Company	**Transworld X-Ray Corp.**
Location	Charlotte, NC, US
Contact	J. A PRINCEHORN
510(k) history	7 submissions · 7 cleared · 1982-1989