FDA 510(k) Premarket Notification

K872468 JOVI MYELOGRAPHY HARNESS

CLEARED

Jul 1, 1987
8 days to decision

K872468 · Product code: **KPR** · Radiology
Source: https://www.510kdatabase.net/k872468/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Jun 23, 1987
Decision date	Jul 1, 1987
Days to decision	8 days
Third-party review	No

APPLICANT

Company	**Jovi, Inc.**
Location	Dallas, TX, US
Contact	DENNIS T GRIGGS
510(k) history	1 submissions · 1 cleared · 1987-1987