FDA 510(k) Premarket Notification

K812332 KERMATH L/C ARM

CLEARED

Sep 16, 1981
29 days to decision

K812332 · Product code: **KPR** · Radiology
Source: https://www.510kdatabase.net/k812332/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Aug 18, 1981
Decision date	Sep 16, 1981
Days to decision	29 days
Third-party review	No

APPLICANT

Company	**Kermath Mfg. Corp.**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1977-1987