FDA 510(k) Premarket Notification

K833559 ARMBOARD

CLEARED

Jan 3, 1984
90 days to decision

K833559 · Product code: **KPR** · Radiology
Source: https://www.510kdatabase.net/k833559/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Oct 5, 1983
Decision date	Jan 3, 1984
Days to decision	90 days
Third-party review	No

APPLICANT

Company	**Mtd, Inc.**
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1977-2007