

K833559 ARMBBOARDJan 3, 1984
90 days to decisionK833559 · 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

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Device record: <https://www.510kdatabase.net/k833559/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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