FDA 510(k) Premarket Notification

K790536 CLEARVIEW

CLEARED

Mar 28, 1979
6 days to decision

K790536 · Product code: **KXJ** · Radiology
Source: https://www.510kdatabase.net/k790536/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Table, Radiologic (KXJ)
Date received	Mar 22, 1979
Decision date	Mar 28, 1979
Days to decision	6 days
Third-party review	No

APPLICANT

Company	**Hipoint Research, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1979-1979