

K790536 CLEARVIEWMar 28, 1979
6 days to decisionK790536 · 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

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Device record: <https://www.510kdatabase.net/k790536/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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