

K840463 DIGITAL CRADLEFeb 21, 1984
19 days to decisionK840463 · Product code: **KXH** · Radiology
Source: <https://www.510kdatabase.net/k840463/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cradle, Patient, Radiologic (KXH)
Date received	Feb 2, 1984
Decision date	Feb 21, 1984
Days to decision	19 days
Third-party review	No

APPLICANT

Company	Spectrum X-Ray Corp.
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1979-1993

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Device record: <https://www.510kdatabase.net/k840463/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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