FDA 510(k) Premarket Notification

K840463 DIGITAL CRADLE

CLEARED

Feb 21, 1984
19 days to decision

K840463 · 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