

K160723 OnSight 3D Extremity SystemAug 30, 2016
167 days to decisionK160723 · Product code: **JAK** · Radiology
Source: <https://www.510kdatabase.net/k160723/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Tomography, Computed (JAK)
Date received	Mar 16, 2016
Decision date	Aug 30, 2016
Days to decision	167 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Carestream Health, Inc.
Location	Rochester, NY, US
Contact	Carolyn Wagner
Website	http://www.carestream.com/default.aspx?LangType=1033
510(k) history	48 submissions · 48 cleared · 2008-2025

Carestream Health, Inc. is a worldwide provider of medical imaging systems and X-ray imaging solutions. The company operates with a manufacturing facility in Rochester, US and maintains a global service and support network across multiple markets. Carestream has received FDA 510(k) clearances from total submissions since 2008. The company specializes in Radiology devices, which represent 94% of its regulatory submissions. The latest clearance was granted in 2025, demonstrating continued active development and market engagement. Recent cleared devices include digital radio...

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