

**K133730 CO PILOT/REGIUS UNITEA**Jan 31, 2014  
56 days to decisionK133730 · Product code: **LLZ** · Radiology  
Source: <https://www.510kdatabase.net/k133730/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Dec 6, 2013
Decision date	Jan 31, 2014
Days to decision	56 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Konica Minolta, Inc.</b>
Location	New York, NY, US
Contact	RUSSELL MUNVES
Website	<a href="http://www.konicaminolta.com">http://www.konicaminolta.com</a>
510(k) history	25 submissions · 25 cleared · 2014-2025

Konica Minolta, Inc. is a global imaging and technology company with a manufacturing facility in New York, US. The company develops advanced diagnostic and imaging solutions for healthcare and industrial applications. Konica Minolta has received FDA 510(k) clearances from total submissions, with no denied submissions on record. The company specializes exclusively in Radiology devices, establishing a focused regulatory portfolio. Its FDA 510(k) clearance history spans from 2014 to 2025, with recent clearances demonstrating continued active development and market engagement...

---