

**K123812 COHERENCE RT THERAPIST CONNECT
 WORKSPACE, V2.3, ARIA CONNECTIVITY OPTION FOR
 SIEMENS NON-IMAGING SYSTEMS, ARIA CONNECTIV**

Mar 1, 2013
 80 days to decision

K123812 · Product code: **IYE** · Radiology
 Source: <https://www.510kdatabase.net/k123812/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accelerator, Linear, Medical (IYE)
Date received	Dec 11, 2012
Decision date	Mar 1, 2013
Days to decision	80 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Siemens Medical Solutions USA, Inc.
Location	Hoffman Estates, IL, US
Contact	CHRISTINE DUNBAR
510(k) history	778 submissions · 778 cleared · 1980-2026

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k123812/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 15, 2026