

K210055 Cios AlphaFeb 5, 2021
28 days to decisionK210055 · Product code: **OWB** · Radiology
Source: <https://www.510kdatabase.net/k210055/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Interventional Fluoroscopic X-ray System (OWB)
Date received	Jan 8, 2021
Decision date	Feb 5, 2021
Days to decision	28 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k210055/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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