

K211713 AngioCloud ServiceApr 28, 2022
329 days to decisionK211713 · Product code: **LLZ** · Radiology
Source: <https://www.510kdatabase.net/k211713/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Jun 3, 2021
Decision date	Apr 28, 2022
Days to decision	329 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Angiocloud, LLC
Location	Atlanta, GA, US
Contact	James Lewis
510(k) history	1 submissions · 1 cleared · 2022-2022

REGULATORY CONSULTANT

Consulting firm	Lighthouse Regulatory Consulting Group, LLC
Contact	Bahram Parvinian

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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