

K260980 cmAngio® V2.0

Jul 30, 2026
127 days to decision

K260980 · Product code: QIH · Radiology
Source: <https://www.510kdatabase.net/k260980/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated Radiological Image Processing Software (QIH)
Date received	Mar 25, 2026
Decision date	Jul 30, 2026
Days to decision	127 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Curemetrix, Inc.
Location	La Jolla, CA, US
Contact	Kevin Harris
510(k) history	4 submissions · 4 cleared · 2019-2026

510k Database - www.510kdatabase.net
Device record: <https://www.510kdatabase.net/k260980/> 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 August 12, 2026