FDA 510(k) Premarket Notification

K260980 cmAngio® V2.0

CLEARED

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