

**K231709 AcuityDR 1013 G4, AcuityDR 1417 G4, AcuityDR 1717G4**Jul 12, 2023  
30 days to decisionK231709 · Product code: **MQB** · Radiology  
Source: <https://www.510kdatabase.net/k231709/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Solid State X-ray Imager (flat Panel/digital Imager) (MQB)
Date received	Jun 12, 2023
Decision date	Jul 12, 2023
Days to decision	30 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Radmedix, LLC</b>
Location	Dayton, OH, US
Contact	Gabriel Issa
510(k) history	6 submissions · 6 cleared · 2020-2023

**REGULATORY CONSULTANT**

---

Consulting firm	<b>Kamm &amp; Associates</b>
Contact	Daniel Kamm

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - [www.510kdatabase.net](https://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k231709/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 25, 2026