

**K251072 Segmentron Viewer**Sep 9, 2025  
155 days to decisionK251072 · Product code: **QIH** · Radiology  
Source: <https://www.510kdatabase.net/k251072/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated Radiological Image Processing Software (QIH)
Date received	Apr 7, 2025
Decision date	Sep 9, 2025
Days to decision	155 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>DGNCT, LLC</b>
Location	Miami, FL, US
Contact	Anastasya Melnikov
510(k) history	2 submissions · 2 cleared · 2025-2026

**REGULATORY CONSULTANT**

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Consulting firm	<b>Hogan Lovells US LLP</b>
Contact	Kelliann Payne

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

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k251072/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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