

**K201272 Geneva**Jul 16, 2020  
65 days to decisionK201272 · Product code: **JAQ** · Radiology  
Source: <https://www.510kdatabase.net/k201272/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Applicator, Radionuclide, Remote-controlled (JAQ)
Date received	May 12, 2020
Decision date	Jul 16, 2020
Days to decision	65 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Nucletron B.V.</b>
Location	Rice Lake, WI, US
Contact	Rigo Meens
510(k) history	11 submissions · 11 cleared · 2013-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k201272/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 26, 2026