

**K051664 SCAN300FP / RETROPAN (VDC)**Jul 20, 2005  
28 days to decisionK051664 · Product code: **MUH** · Radiology  
Source: <https://www.510kdatabase.net/k051664/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Extraoral Source, Digital (MUH)
Date received	Jun 22, 2005
Decision date	Jul 20, 2005
Days to decision	28 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Oy Ajat, Ltd.</b>
Location	Ormond Beach, FL, US
Contact	CLAUDE D BERTHOIN
510(k) history	1 submissions · 1 cleared · 2005-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k051664/> 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 18, 2026