

**K180663 LaVid FMTS DIAGNOSTIC X-RAY SYSTEM**May 18, 2018  
65 days to decisionK180663 · Product code: **KPR** · Radiology  
Source: <https://www.510kdatabase.net/k180663/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Mar 14, 2018
Decision date	May 18, 2018
Days to decision	65 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Livemoretech, Inc.</b>
Location	Plano, TX, US
Contact	Jae Hong Kim
510(k) history	6 submissions · 6 cleared · 2018-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k180663/> 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 27, 2026