

K163624 Dynamic Visualization II Image Processing Option for the FDX Console (DR-ID300CL of the DR-ID600)

Jun 30, 2017
190 days to decision

K163624 · Product code: **MQB** · Radiology
Source: <https://www.510kdatabase.net/k163624/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Solid State X-ray Imager (flat Panel/digital Imager) (MQB)
Date received	Dec 22, 2016
Decision date	Jun 30, 2017
Days to decision	190 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Fujifilm Medical Systems U.S.A, Inc.
Location	Stamford, CT, US
Contact	Dharaben Desai
510(k) history	39 submissions · 39 cleared · 2005-2018

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k163624/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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