FDA 510(k) Premarket Notification

K180663 LaVid FMTS DIAGNOSTIC X-RAY SYSTEM

CLEARED

May 18, 2018
65 days to decision

K180663 · Product code: **KPR** · Radiology
Source: https://www.510kdatabase.net/k180663/

SUBMISSION DETAILS

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

Company	**Livermoretech, Inc.**
Location	Plano, TX, US
Contact	Jae Hong Kim
510(k) history	6 submissions · 6 cleared · 2018-2025