FDA 510(k) Premarket Notification

K193535 EZER, Portable X-ray System

CLEARED

May 7, 2020
139 days to decision

K193535 · Product code: **IZL** · Radiology
Source: https://www.510kdatabase.net/k193535/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Mobile (IZL)
Date received	Dec 20, 2019
Decision date	May 7, 2020
Days to decision	139 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k193535/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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