FDA 510(k) Premarket Notification

K200990 VIDAvision

CLEARED

Aug 7, 2020
114 days to decision

K200990 · Product code: **JAK** · Radiology
Source: https://www.510kdatabase.net/k200990/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Tomography, Computed (JAK)
Date received	Apr 15, 2020
Decision date	Aug 7, 2020
Days to decision	114 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Vida Diagnostics, Inc.**
Location	Iowa City, IA, US
Contact	Sandra Stapleton
510(k) history	2 submissions · 2 cleared · 2008-2020