FDA 510(k) Premarket Notification

K173425 SRT-100+

CLEARED

Mar 23, 2018
141 days to decision

K173425 · Product code: **JAD** · Radiology
Source: https://www.510kdatabase.net/k173425/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Therapeutic, X-ray (JAD)
Date received	Nov 2, 2017
Decision date	Mar 23, 2018
Days to decision	141 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Sensus Healthcare, Inc.**
Location	Boca Raton, FL, US
Contact	Nicolas Soro
510(k) history	4 submissions · 4 cleared · 2018-2019

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k173425/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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