FDA 510(k) Premarket Notification

K220663 AccuCTP

CLEARED

Nov 22, 2022
260 days to decision

K220663 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k220663/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Mar 7, 2022
Decision date	Nov 22, 2022
Days to decision	260 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**ArteryFlow Technology Co., Ltd.**
Location	Hangzhou City, CN
Contact	Jianping Xiang
510(k) history	6 submissions · 6 cleared · 2021-2023