FDA 510(k) Premarket Notification

K220256 MIM-Ablation

CLEARED

Oct 7, 2022
249 days to decision

K220256 · Product code: **QTZ** · Radiology
Source: https://www.510kdatabase.net/k220256/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Radiological Image Processing Software For Ablation Therapy Planning And Evaluation (QTZ)
Date received	Jan 31, 2022
Decision date	Oct 7, 2022
Days to decision	249 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Mim Software, Inc.**
Location	Cleveland, OH, US
Contact	Daniel Darkow
510(k) history	22 submissions · 22 cleared · 2011-2026