FDA 510(k) Premarket Notification

K013481 DUAL ENERGY AND TISSUE EQUALIZATION SOFTWARE OPTION

CLEARED

Nov 2, 2001
14 days to decision

K013481 · Product code: **KPR** · Radiology
Source: https://www.510kdatabase.net/k013481/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Oct 19, 2001
Decision date	Nov 2, 2001
Days to decision	14 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**GE Medical Systems**
Location	Milwaukee, WI, US
Contact	JODI PARKER
510(k) history	169 submissions · 166 cleared · 1989-2022