

**K013481 DUAL ENERGY AND TISSUE EQUALIZATION
SOFTWARE OPTION**Nov 2, 2001
14 days to decisionK013481 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k013481/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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