FDA 510(k) Premarket Notification

CLEARED

K080582 MEDICATECH, MODEL DDR MAK-800, DDT MAK-1000 FS, DDR MAK-1100 FA

May 15, 2008
73 days to decision

K080582 · Product code: **KPR** · Radiology
Source: https://www.510kdatabase.net/k080582/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Mar 3, 2008
Decision date	May 15, 2008
Days to decision	73 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Medicatech USA**
Location	Deer Field, IL, US
Contact	DANIEL KAMM
510(k) history	9 submissions · 9 cleared · 2008-2023