

**K080582 MEDICATECH, MODEL DDR MAK-800, DDT MAK-1000  
FS, DDR MAK-1100 FA**May 15, 2008  
73 days to decisionK080582 · Product code: **KPR** · Radiology  
Source: <https://www.510kdatabase.net/k080582/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k080582/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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