

K011111 MXD100/MX 20, OR MX 30, OR MX 40 OR MX 50--(MXD 100 SERIES)May 30, 2001
49 days to decisionK011111 · Product code: **KPR** · Radiology
Source: <https://www.510kdatabase.net/k011111/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Stationary (KPR)
Date received	Apr 11, 2001
Decision date	May 30, 2001
Days to decision	49 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Modular X-Ray Devices
Location	Roodepoort, ZA
Contact	S.B.B. DESEMBERG
510(k) history	1 submissions · 1 cleared · 2001-2001

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k011111/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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