

K082347 DIGITAL RADIOGRAPHY, FLEXDR C30Nov 18, 2008
95 days to decisionK082347 · Product code: **MQB** · Radiology
Source: <https://www.510kdatabase.net/k082347/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Solid State X-ray Imager (flat Panel/digital Imager) (MQB)
Date received	Aug 15, 2008
Decision date	Nov 18, 2008
Days to decision	95 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Konica Minolta Technoproduct Co., Ltd.
Location	New York, NY, US
Contact	RUSSEL MUNVES
510(k) history	1 submissions · 1 cleared · 2008-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k082347/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 17, 2026