

K103703 REGUIS SIGMAJun 2, 2011
164 days to decisionK103703 · Product code: **MQB** · Radiology
Source: <https://www.510kdatabase.net/k103703/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Solid State X-ray Imager (flat Panel/digital Imager) (MQB)
Date received	Dec 20, 2010
Decision date	Jun 2, 2011
Days to decision	164 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Konica Minolta Medical & Graphic, Inc.
Location	Hachioji-Shi Tokyo, JP
Contact	RUSSELL MUNVES
510(k) history	27 submissions · 27 cleared · 2002-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k103703/> 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