

K120263 VOLUX 21C, DIGITAL EXTRAORAL SOURCE X-RAY SYSTEM

Jul 10, 2012
162 days to decision

K120263 · Product code: **MUH** · Radiology
Source: <https://www.510kdatabase.net/k120263/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, X-ray, Extraoral Source, Digital (MUH)
Date received	Jan 30, 2012
Decision date	Jul 10, 2012
Days to decision	162 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Genoray Co., Ltd.
Location	Flintville, TN, US
Contact	JAE KIM
510(k) history	24 submissions · 24 cleared · 2007-2026

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k120263/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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