FDA 510(k) Premarket Notification

K103038 DUET DRF

CLEARED

Mar 29, 2011
166 days to decision

K103038 · Product code: **JAA** · Radiology
Source: https://www.510kdatabase.net/k103038/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	System, X-ray, Fluoroscopic, Image-intensified (JAA)
Date received	Oct 14, 2010
Decision date	Mar 29, 2011
Days to decision	166 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cmt Medical Technologies, Ltd.**
Location	Haifa, IL
Contact	SHLOMI DINES
510(k) history	12 submissions · 12 cleared · 1995-2020

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k103038/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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