FDA 510(k) Premarket Notification

K133320 SICAT FUNCTION

CLEARED

Feb 25, 2014
120 days to decision

K133320 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k133320/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Oct 28, 2013
Decision date	Feb 25, 2014
Days to decision	120 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Sicat GmbH & Co. KG**
Location	Bonn, DE
Contact	MANFRED BREUER
510(k) history	9 submissions · 9 cleared · 2006-2019

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k133320/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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