FDA 510(k) Premarket Notification

CLEARED

K053010 VISTADENT AT COMPLETE, MODEL 3.1

Nov 22, 2005
27 days to decision

K053010 · Product code: **LLZ** · Radiology
Source: https://www.510kdatabase.net/k053010/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Image Processing, Radiological (LLZ)
Date received	Oct 26, 2005
Decision date	Nov 22, 2005
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Dentsply Intl., Inc.**
Location	York, PA, US
Contact	HELEN LEWIS
510(k) history	12 submissions · 12 cleared · 2005-2014