FDA 510(k) Premarket Notification

K150585 Ascent 3D

CLEARED

Dec 31, 2015
297 days to decision

K150585 · Product code: **EBZ** · Dental
Source: https://www.510kdatabase.net/k150585/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Activator, Ultraviolet, For Polymerization (EBZ)
Date received	Mar 9, 2015
Decision date	Dec 31, 2015
Days to decision	297 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**CAO Group, Inc.**
Location	West Jordan, UT, US
Contact	Robert K Larsen
Website	https://www.caogroup.com
510(k) history	31 submissions · 31 cleared · 2001-2026

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k150585/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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