

**K072643 CRYSTALEYE SPECTROPHOTOMETER/CRYSTALEYE
BASIC SET, MODEL CE100-DC/US**Nov 16, 2007
59 days to decisionK072643 · Product code: **KZN** · Dental
Source: <https://www.510kdatabase.net/k072643/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Scanner, Color (KZN)
Date received	Sep 18, 2007
Decision date	Nov 16, 2007
Days to decision	59 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Olympus Corporation
Location	Melville, NY, US
Contact	FUMIAKI KANAI
Website	http://www.olympus-global.com/en/global
510(k) history	5 submissions · 5 cleared · 2004-2007

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k072643/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 18, 2026