

**K072572 PRIMACONNEX RD AND WD ESTHETIC CONTOUR ZI
ABUTMENTS**Oct 5, 2007
23 days to decisionK072572 · Product code: **NHA** · Dental
Source: <https://www.510kdatabase.net/k072572/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Abutment, Implant, Dental, Endosseous (NHA)
Date received	Sep 12, 2007
Decision date	Oct 5, 2007
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Lifecore Biomedical, Inc.
Location	Washington, DC, US
Contact	KAREN CLEMENT
Website	http://www.lifecore.com/
510(k) history	34 submissions · 34 cleared · 1991-2007

Lifecore Biomedical, Inc. is a sterile injectable contract development and manufacturing organization (CDMO) and sodium hyaluronate producer with a manufacturing facility in Washington, US. The company specializes in formulation development, aseptic fill-finish, analytical testing, and stability services for pharmaceutical and medical device manufacturers. Lifecore received FDA 510(k) clearances from total submissions, with clearances spanning 1991 to 2007. The company's regulatory focus was predominantly Dental devices, representing 97% of submissions. This includes dent...

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