

K970776 LIFECORE O-RING ABUTMENTMay 27, 1997
84 days to decisionK970776 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k970776/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Mar 4, 1997
Decision date	May 27, 1997
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary
Other names	LIFECORE DALLA BONA ABUTMENT

APPLICANT

Company	Lifecore Biomedical, Inc.
Location	Washington, DC, US
Contact	DONNA BAHLS
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...
