

**K944068 RESTORE SELF TAPPING DENTAL IMPLANT SYST (5, 5.5,6MM)**Feb 2, 1995  
171 days to decisionK944068 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k944068/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Aug 15, 1994
Decision date	Feb 2, 1995
Days to decision	171 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Lifecore Biomedical, Inc.</b>
Location	Washington, DC, US
Contact	LYNN CUPERUS
Website	<a href="http://www.lifecore.com/">http://www.lifecore.com/</a>
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...

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k944068/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 26, 2026