

**K052070 ATLANTIS ABUTMENT IN ZIRCONIA, ATLANTIS GEMINI ABUTMENT IN ZIRCONIA, ATLANTIS GEMEINI + ABUTMENT IN ZIRCONIA**Oct 14, 2005  
74 days to decisionK052070 · Product code: **NHA** · Dental  
Source: <https://www.510kdatabase.net/k052070/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Abutment, Implant, Dental, Endosseous (NHA)
Date received	Aug 1, 2005
Decision date	Oct 14, 2005
Days to decision	74 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Atlantis Components, Inc.</b>
Location	Skokie, IL, US
Contact	BETSY A BROWN
510(k) history	18 submissions · 18 cleared · 1999-2011

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k052070/>. 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 20, 2026